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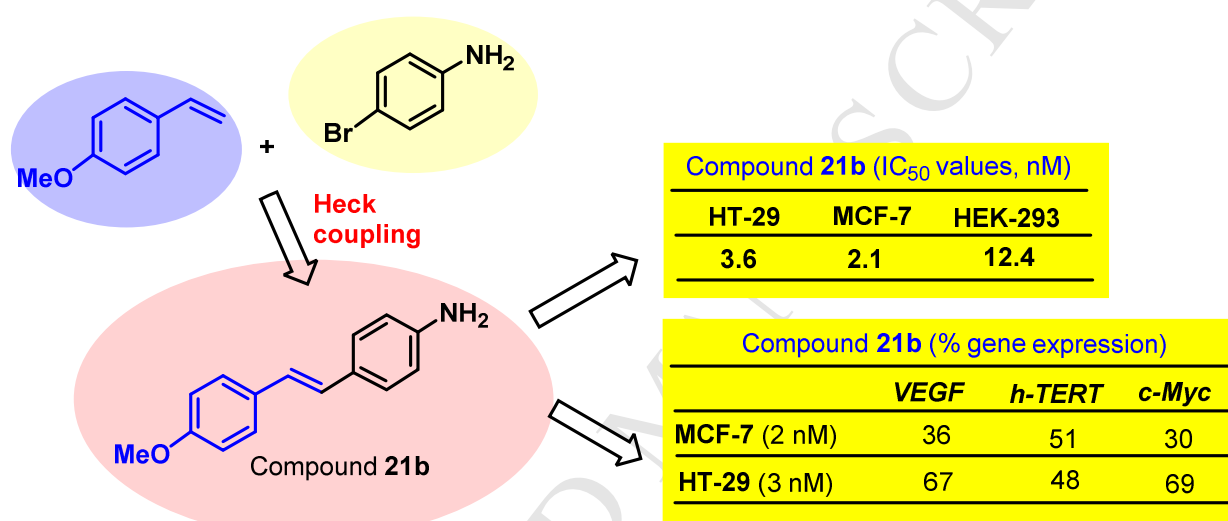
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Inhibitory effect of cytotoxic stilbenes related to resveratrol on the expression of the VEGF, hTERT and c-Myc genes

Rosa Martí-Centelles, Eva Falomir [*] *Juan Murga, Miguel Carda* [*] *and J. Alberto Marco*

Graphical Abstract

Thirty-nine stilbenes have been prepared. Their cytotoxicities and ability to inhibit the expression of VEGF, h-TERT and c-Myc genes have been measured. One compound is very active in these properties.

Inhibitory effect of cytotoxic stilbenes related to resveratrol on the expression of the VEGF, hTERT and c-Myc genes

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Highlights

- Thirty-nine stilbene derivatives related to resveratrol were prepared through Heck type coupling reactions

- The molecules were tested for cytotoxicity on a normal and two tumoral cell lines

- The molecules were tested as regards inhibition of secretion of VEGF protein

- The molecules were tested as regards downregulation of *VEGF*, *h-TERT* and *c-Myc* genes

- (*E*)-4-(4-methoxystyryl)aniline showed cytotoxicity at the low nanomolar level and was able to significantly inhibit secretion of VEGF protein and the expression of the *VEGF*, *h-TERT* and *c-Myc* genes at similarly low concentrations

Inhibitory effect of cytotoxic stilbenes related to resveratrol on the expression of the VEGF, hTERT and c-Myc genes

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ABSTRACT

A group of thirty-nine stilbene derivatives, prepared by means of Heck coupling reactions, has been investigated for their cytotoxicity, as well as for their ability to inhibit the production of the vascular endothelial growth factor (VEGF) and the activation of telomerase. The ability of these compounds to inhibit proliferation of two tumoral cell lines (HT-29 and MCF-7) and one non tumoral cell line (HEK-293) was first determined. Subsequently, we determined the capacity of the compounds to inhibit the secretion of VEGF in the aforementioned cell lines and to downregulate the expression of the *VEGF*, *h-TERT* and *c-Myc* genes, the two latter involved in the control of the activation of telomerase. One of the synthetic stilbenes, (*E*)-4-(4-methoxystyryl)aniline, showed strong cytotoxicity and proved able to cause a marked decrease both in the secretion of VEGF and in the expression of the *h-TERT* and *c-Myc* genes, in all cases at concentrations in the low nanomolar range.

KEYWORDS

resveratrol, stilbenes, VEGF, gene regulation, telomerase, *hTERT*, *c-Myc*.

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1. Introduction

Resveratrol, a natural compound belonging to the class of stilbenes (Fig. 1) is a polyphenolic compound which occurs in grapes, red wine, chocolate, and certain berries and roots. It became popular since the 1993 paper on its inhibitory effects against the oxidation of human low-density lipoproteins, a chemical event at the onset of the pathogenesis of atherosclerosis [1]. In later years, further important biological features were reported for the compound, most particularly antioxidant, anticarcinogenic, and anti-inflammatory properties [2]. These properties were related to the ability of the compound to retard the onset of age-related diseases [3] and to its beneficial effect on other ailments such as viral attacks [4], diabetes [5], inflammation [6] and neurodegenerative diseases [7]. In addition, resveratrol has been shown to display, among other features: a) cancer chemopreventive effects by means of inhibiting the initiation step of the carcinogenesis process [8]; b) anticancer activity through inhibition of cell cycle progression and induction of apoptotic cell death [9]; and c) antiangiogenic activity with sufficient potency to suppress FGF-2 and VEGF-induced neovascularisation in vivo [10] and to inhibit bovine aorta endothelial cell proliferation, migration and tube formation in vitro [11].
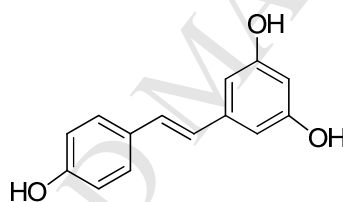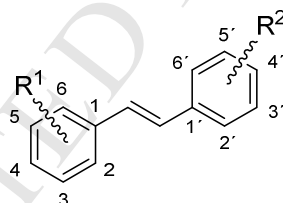

Figure 1. Structure of resveratrol

Tumor angiogenesis is a very complex process and involves the tight interplay of many factors [12]. Vascular endothelial growth factor (VEGF) is an inducer of angiogenesis and promotes endothelial cell survival, proliferation and migration while increasing vascular permeability [13]. In fact, overexpression in the production of VEGF has been reported to occur in various types of tumors [14]. Not unexpectedly, VEGF has become one further key target molecule in cancer therapy [15,16].

Cancer cells are able to evolve the ability to overcome senescence upon telomerase reactivation [17]. This enzyme has been detected in about 90% of all malignant tumors [18], so that drugs with ability to inhibit telomerase activity are potentially useful weapons in the fight against cancer, aging and other diseases related to premature telomere shortening [19]. Recently, it has been reported that resveratrol downregulates the telomerase activity of MCF-7 cells and the nuclear levels of hTERT, the reverse transcriptase subunit of the telomerase complex [20].

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2. Research purpose

When aiming at a more efficient anticancer therapy, compounds exhibiting not only high cytotoxic activity but also the ability to suppress tumoral angiogenesis and the expression of genes related to telomerase activation would be highly desirable [21]. On our ongoing research on anticancer properties of synthetic analogues derived from natural products, we have recently reported on the inhibition of VEGF formation in cancer cells and on endothelial cell differentiation promoted by synthetic stilbene derivatives [22]. Our study has established that resveratrol and several stilbene derivatives lead to a decrease in the production of VEGF by HT-29 cells. In view of these precedents we decided to evaluate the influence of new synthetic stilbene analogues not only on VEGF production, but also on the expression of the *hTERT* gene, involved in the process of telomerase activation. In addition, we further decided to measure the inhibition of expression of *c-Myc*, an oncogene which encodes the c-Myc protein that acts as a transcription factor for *hTERT* expression.

The structures of the synthetic resveratrol analogues which are the subject of this work are depicted in Figure 2. These compounds belong to two types: one corresponds to stilbenes carrying hydroxy and/or methoxy groups (*a*-series in Figure 2), whereas the other (*b*-series) includes stilbenes carrying amine or amide functions in one of the rings, as well as hydroxyl or methoxy groups in the other.


1a R^1 = 2-OH R^2 = 2´-OH	**6a** R^1 = 4-OH R^2 = 4´-OH	**11a** R^1 = 3-OMe R^2 = 3´-OH
2a R^1 = 3-OH R^2 = 2´-OH	**7a** R^1 = 2-OMe R^2 = 2´-OH	**12a** R^1 = 4-OMe R^2 = 3´-OH
3a R^1 = 4-OH R^2 = 2´-OH	**8a** R^1 = 3-OMe R^2 = 2´-OH	**13a** R^1 = 2-OMe R^2 = 4´-OH
4a R^1 = 3-OH R^2 = 3´-OH	**9a** R^1 = 4-OMe R^2 = 2´-OH	**14a** R^1 = 3-OMe R^2 = 4´-OH
5a R^1 = 4-OH R^2 = 3´-OH	**10a** R^1 = 2-OMe R^2 = 3´-OH	**15a** R^1 = 4-OMe R^2 = 4´-OH
1b R^1 = 2-NH$_2$ R^2 = H	**9b** R^1 = 3-NH$_2$ R^2 = 4´-OH	**17b** R^1 = 3-NH$_2$ R^2 = 3´-OMe
2b R^1 = 3-NH$_2$ R^2 = H	**10b** R^1 = 4-NH$_2$ R^2 = 2´-OH	**18b** R^1 = 3-NH$_2$ R^2 = 4´-OMe
3b R^1 = 4-NH$_2$ R^2 = H	**11b** R^1 = 4-NH$_2$ R^2 = 3´-OH	**19b** R^1 = 4-NH$_2$ R^2 = 2´-OMe
4b R^1 = 2-NH$_2$ R^2 = 2´-OH	**12b** R^1 = 4-NH$_2$ R^2 = 4´-OH	**20b** R^1 = 4-NH$_2$ R^2 = 3´-OMe
5b R^1 = 2-NH$_2$ R^2 = 3´-OH	**13b** R^1 = 2-NH$_2$ R^2 = 2´-OMe	**21b** R^1 = 4-NH$_2$ R^2 = 4´-OMe
6b R^1 = 2-NH$_2$ R^2 = 4´-OH	**14b** R^1 = 2-NH$_2$ R^2 = 3´-OMe	**22b** R^1 = 2-NHCO(CH$_2$)$_{10}$Me R^2 = H
7b R^1 = 3-NH$_2$ R^2 = 2´-OH	**15b** R^1 = 2-NH$_2$ R^2 = 4´-OMe	**23b** R^1 = 3-NHCO(CH$_2$)$_{10}$Me R^2 = H
8b R^1 = 3-NH$_2$ R^2 = 3´-OH	**16b** R^1 = 3-NH$_2$ R^2 = 2´-OMe	**24b** R^1 = 4-NHCO(CH$_2$)$_{10}$Me R^2 = H

Figure 2. Structures of synthetic resveratrol analogues.

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3. Synthetic work

Stilbene derivatives **1a**-**15a** were prepared by means of palladium-catalyzed Heck-type cross coupling reactions [23]. Monohydroxylated styrenes were used as the nucleophilic partners whereas hydroxy or methoxy bromobenzenes played the role of the electrophilic components, with $Pd(PPh_3)_4$ being the catalyst [24] (conditions A in Scheme 1). These reactions were performed under microwave irradiation, as these conditions have often been found to accelerate this kind of coupling reactions [25] (for details and yields, see Experimental section).
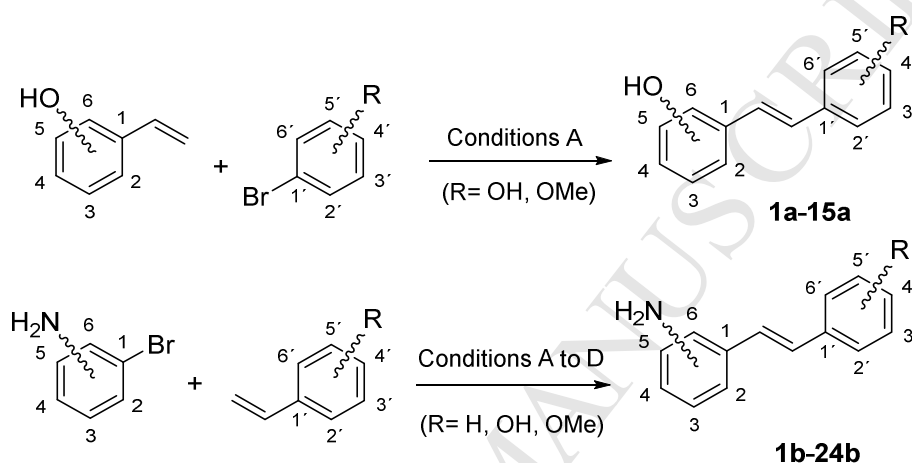
Scheme 1. Synthesis of stilbene derivatives by means of palladium-catalyzed Heck couplings (for more detailed descriptions of the reaction conditions and yields, see Supporting Information).

Preparation of compounds belonging to the *b*-series was also carried out with Heck-type cross coupling reactions. The same type of nucleophilic partners as above were used, with the electrophilic components being now bromoanilines. The aforementioned reaction conditions A, however, used in the synthesis of compounds of the *a*-series, were successful in the preparation of some compounds (e. g. **4b**, **5b**, **7b**, **9b**), but proved inefficient in other cases or else failed completely. In order to obtain the remaining amino stilbenes without protection of the amino groups, we undertook a survey of catalysts and conditions which resulted in three new reactions conditions, B, C and D (Scheme 1). In this case, $Pd(NH_3)_2Cl_2$ was the catalyst [26], the differences residing in the reaction conditions. All Heck reactions were highly stereoselective and afforded, after chromatographic purification, the corresponding stilbenes of *E*-configuration with no traces of *Z*-isomers. Yields ranged from 50% to 88% and, most importantly, no protection/deprotection steps were needed.

Amides **22b**-**24b** were prepared with the aim at observing whether the introduction of a lipophilic side chain would cause visible changes in the biological properties. Amide **23b** was synthesized upon

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coupling compound **2b** with dodecanoic acid in the presence of chlorosulfonyl isocyanate [27]. Attempts at preparing **22b** and **24b** in the same way by means of acylation of **1b** and **3b**, respectively, were unsatisfactory. Thus, **22b** and **24b** were obtained by first acylating the corresponding bromoaniline followed by Heck coupling of the resulting amide with styrene under conditions C.

4. Biological work

4.1. Inhibition of cell proliferation

We carried out a measurement of the cytotoxic activity of our synthetic stilbenes using two tumoral cells, the human colon HT-29 and the breast adenocarcinoma MCF-7 cell lines, as well as one non tumoral cell line, the human embryonic kidney cell line, HEK-293 [28]. Table 1 shows cytotoxicity values for stilbenes of the *a*-series, expressed as the compound concentration (µg/mL) that causes 50% inhibition of cell growth (IC_{50}). Table 1 further shows the α and β coefficients, obtained by dividing the IC_{50} values of the normal cell line by those of either the HT-29 or the MCF-7 cell line, respectively (see footnote in Table 1). The higher the value of either coefficient, the higher the therapeutic safety margin of the compound in the corresponding cell line.

Table 1. IC_{50} values (µg/mL) and selectivity coefficients for compounds of the *a*-series.

Compound	HT-29	MCF-7	HEK-293	α^b	β^c
Resveratrol	34.1 ± 1.4	16.1 ± 1.1	7.1 ± 1.0	0.2	0.4
1a	25 ± 7	23.0 ± 1.3	24.3 ± 2.3	0.9	1.1
2a	27 ± 7	20.1 ± 2.1	18 ± 8	0.7	0.9
3a	22 ± 9	21 ± 3	17.5 ± 1.5	0.8	0.8
4a	25.9 ± 1.0	21 ± 6	21 ± 5	0.8	1
5a	6.9 ± 1.9	23 ± 3	1.1 ± 0.7	0.2	< 0.1
6a	>100	6.8 ± 0.2	1.2 ± 0.1	< 0.1	0.2
7a	23 ± 6	17.9 ± 2.4	15.0 ± 2.5	0.7	0.8
8a	26 ± 3	18 ± 4	20 ± 4	0.8	1.1
9a	10 ± 4	17 ± 6	21 ± 4	2.1	1.2
10a	17 ± 4	24.3 ± 1.0	24 ± 3	1.4	1
11a	11.5 ± 3	13 ± 3	19 ± 7	1.6	1.5
12a	22 ± 4	21 ± 5	24 ± 3	1.1	1.1
13a	24 ± 3	19 ± 4	41 ± 3	1.7	2.2
14a	26 ± 4	21.0 ± 1.5	24 ± 3	0.9	1.1

|					
15a	17.4 ± 0.2	11.3 ± 2.3	3.9 ± 0.5	0.2	0.3

[a]Values are the average ($\pm$ s.d.) of three different measurements performed as described in the Material and Methods section. [b]$\alpha = IC_{50}$ (HEK-293) / IC_{50} (HT-29). [c]$\beta = IC_{50}$ (HEK-293) / IC_{50} (MCF-7). Values of α and β have been rounded off to a decimal figure.

The observed values for IC_{50} are in the low to medium micromolar range, with compound **5a** showing the lowest values in the HT-29 cell line and compound **6a** showing the lowest values in the MCF-7 cell line. In the case of HT-29 cell lines, all synthetic compounds exhibit better IC_{50} values than resveratrol. As regards the α and β coefficients, the calculated values are is most cases close to, or even lower than 1. This means that the cytotoxic activity of these compounds is similar in both the non tumoral and the tumoral line.

Table 2. IC_{50} values (μg/mL) and selectivity coefficients for compounds of the *b*-series.

Compound	HT-29	MCF-7	HEK-293	α[b]	β[c]
Resveratrol	34.1 ± 1.4	16.1 ± 1.1	7.1 ± 1.0	0.2	0.4
1b	17 ± 4	1.4 ± 0.1	1.3 ± 0.2	0.1	0.9
2b	12.9 ± 0.9	14 ± 4	84 ± 11	6.5	6
3b	4.4 ± 1.1	16 ± 0.1	11.4 ± 0.4	2.6	0.7
4b	36 ± 5	39 ± 3	27 ± 7	0.7	0.7
5b	30 ± 5	17 ± 3	20 ± 7	0.7	1.2
6b	>100	79 ± 3	7 ± 3	< 0.1	< 0.1
7b	30.4 ± 1.8	24.7 ± 0.2	96 ± 15	3.2	3.9
8b	43 ± 16	54 ± 7	>100	2.3	1.8
9b	51 ± 16	6 ± 3	7 ± 3	0.1	1.2
10b	37 ± 5	23 ± 4	21.4 ± 1.4	0.6	1
11b	42 ± 6	24.8 ± 0.4	25 ± 8	0.6	1
12b	30.6 ± 2.4	2.2 ± 0.4	0.8 ± 0.1	0.03	0.4
13b	>100	>100	2.1 ± 0.7	< 0.1	< 0.1
14b	21.2 ± 1.5	6.1 ± 1.8	0.5 ± 0.1	< 0.1	< 0.1
15b	73 ± 5	0.4 ± 0.1	0.6 ± 0.1	< 0.1	1.4
16b	25.3 ± 0.1	38.7 ± 0.6	45 ± 0.3	1.8	1.2
17b	19.6 ± 2.0	34 ± 5	16 ± 5	0.8	0.5
18b	16.3 ± 0.4	23 ± 4	11 ± 0.3	0.7	0.5
19b	22 ± 4	3.2 ± 1.7	1 ± 0.3	< 0.1	0.3
20b	16.1 ± 0.1	52 ± 6	18 ± 4	1.1	0.3
21b	0.0036 ± 0.0015	0.0021 ± 0.0005	0.012 ± 0.004	3.4	5.7
22b	2.5 ± 0.7	12 ± 0.9	16 ± 9	6.4	1.3

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23b	>100	3.5 ±0.5	11.6 ± 0.6	< 0.1	3.3
24b	>100	5.8 ± 0.1	10 ± 4	< 0.1	1.7

[a]Values are the average (± s.d.) of three different measurements performed as described in the Material and Methods section. [b]α = IC$_{50}$ (HEK-293) / IC$_{50}$ (HT-29). [c]β = IC$_{50}$ (HEK-293) / IC$_{50}$ (MCF-7). Values of α and β have been rounded off to a decimal figure.

Cytotoxicity values for stilbenes of the *b*-series are shown in Table 2, with most compounds showing cytotoxicities of the same order of magnitude as those of Table 1. It can be further observed that the introduction of the aliphatic side chains (compounds **22b**-**24b**) did not cause improvements in the cytotoxic activities. However, one of the stilbene derivatives calls for attention: compound **21b** showed cytotoxic values in the low nanomolar range for both cancer cell lines. The α and β coefficients, clearly above 1, further show that the compound is noticeably more cytotoxic for tumoral cell lines than for normal ones, a desirable feature.

4.2. Effect of stilbene derivatives on VEGF protein secretion and VEGF gene inhibition

The action of stilbene derivatives on VEGF secretion and *VEGF* gene inhibition was conducted on the HT-29 tumoral cell line. For these measurements compounds **9a**-**13a**, **2b**, **3b**, **7b**, **8b**, **16b** and **20b**-**22b** were selected because they had α-coefficients above 1. In addition, compounds **1b**, **14b**, **17b** and **18b** were further selected because they had lower IC$_{50}$ values than resveratrol.

The secretion of the vascular endothelial growth factor (VEGF) was first determined by means of the ELISA procedure [29], as described in the Experimental Section. Results for all the selected compounds (except **21b**, which will be discussed later) are depicted in Figure 3, which shows the percentage of VEGF secreted to the culture medium after 72 h of incubation in the presence of DMSO (control experiment) and in the presence of each of the compounds investigated at a concentration of 10 μg/mL (lower than their IC$_{50}$ values).
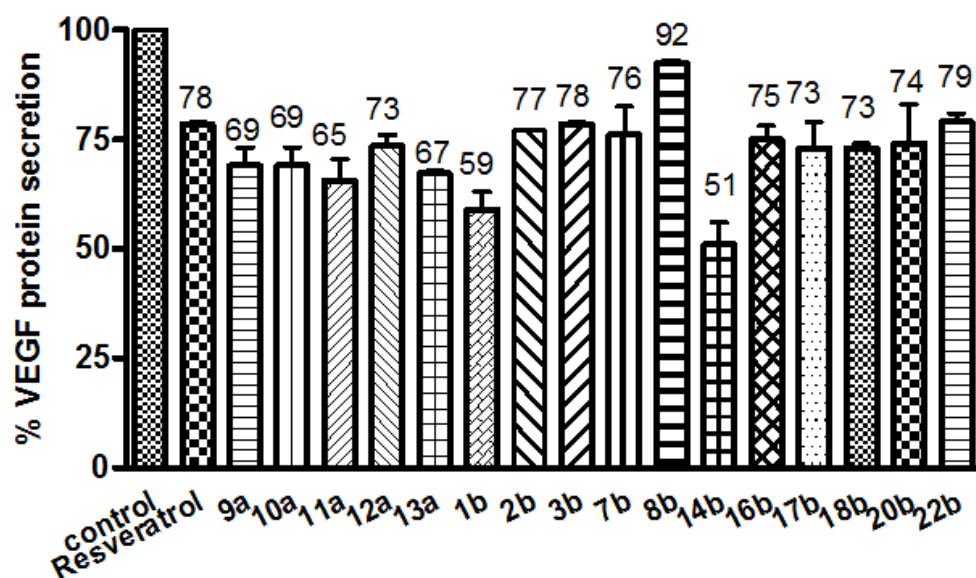
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Figure 3. VEGF protein secretion from HT-29 cells determined by means of the ELISA procedure. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

While the majority of the compounds showed an inhibitory effect on VEGF production, the values were very similar to that of resveratrol, which reduced VEGF secretion to 78% of the control value. The most active is compound **14b**, which causes a decrease to 51%.

The high cytotoxicity of compound **21b**, as compared with the remaining stilbenes, justified a separate study. It was evaluated as regards inhibition of VEGF production on both the HT-29 and the MCF-7 cell line (Figure 4). When a concentration of 1 ng/mL was used, no noticeable effect on VEGF production was detected in either of the two cell lines. However, at 2 ng/mL concentration, compound **21b** inhibited VEGF secretion to 84% of the control value in the case of HT-29 cells and to 75% in the case of MCF-7. When HT-29 cells were cultivated in the presence of 3 ng/mL of compound **21b**, VEGF protein secretion further dropped to 59%. *These values are similar to that achieved by resveratrol at a concentration of 10 µg/mL.* Compound **21b** therefore is three orders of magnitude more active than resveratrol in inhibiting VEGF production.
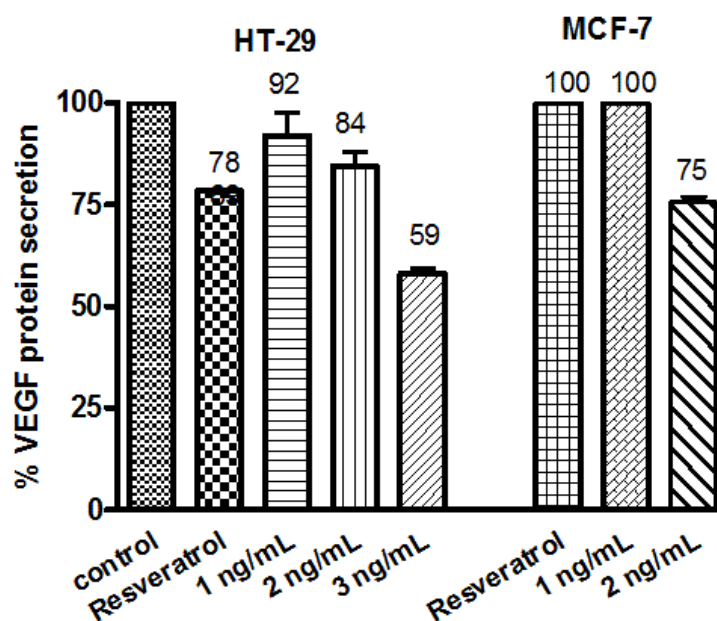
Figure 4. VEGF protein secretion from HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).
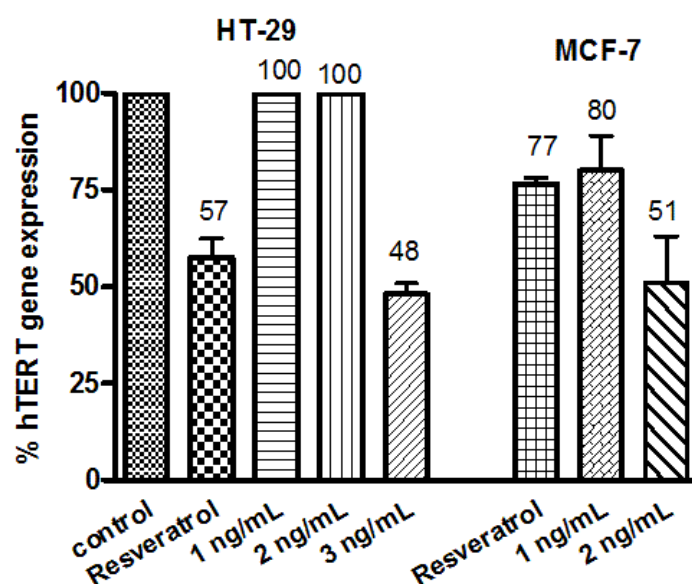
While the results depicted in Figures 3 and 4 point out that the investigated compounds cause inhibition of the VEGF production, they do not permit conclusions about the precise phase of the VEGF generation process with which they interfere. In order to further investigate the mechanism of action of the compounds under study, we proceeded to determine whether they were able to control protein production at the transcriptional level. The VEGF production we have determined with the ELISA procedure is that secreted to the culture medium and corresponds mainly to the lighter VEGFA-165 isoform [30]. Therefore, in order to evaluate the ability of our compounds to inhibit the expression of the *VEGF* gene, we incubated HT-29 cells for 48 h with several selected stilbene derivatives at a concentration of 10 µg/mL, again with DMSO as the control test. For that purpose, the reverse transcription quantitative PCR (RT-qPCR) methodology [31] was used as described in the Experimental Section. Results for all the selected compounds (except **21b**) are shown in Figure 5 as percentage of the *VEGF* gene expression. All values were standardized (100%) to control (DMSO) and to β-actine.
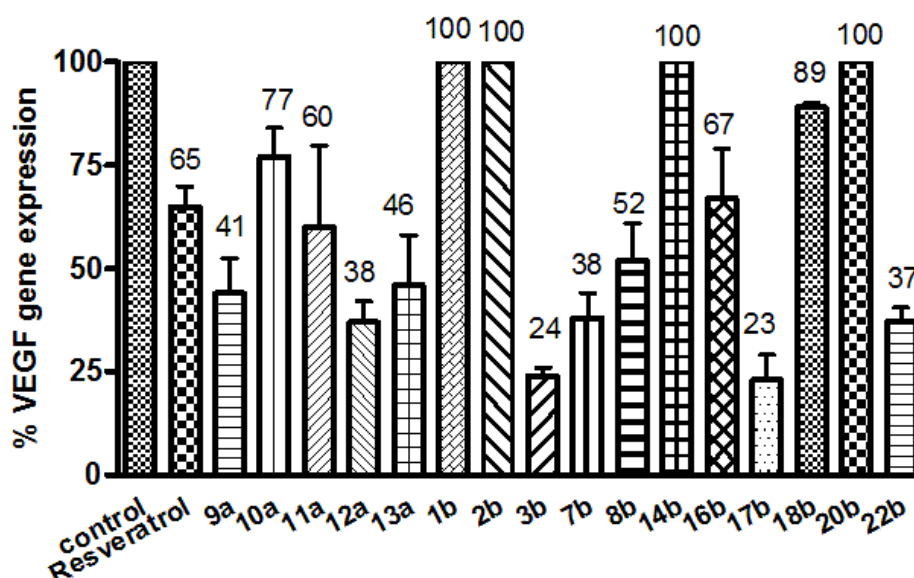
Figure 5. Expression percentage of the *VEGF* gene after 48 h of incubation of HT-29 cells determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).
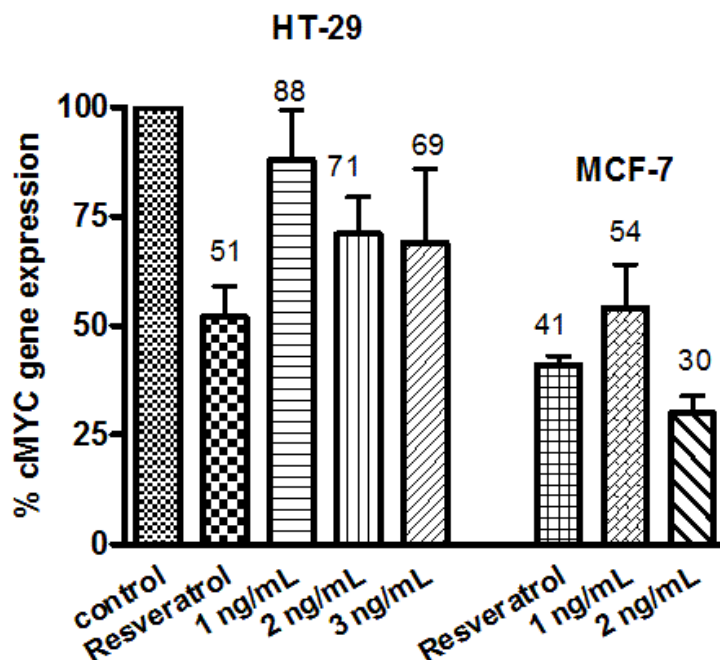
Comparison of Fig. 5 with Fig. 4 reveals much more meaningful differences in activity between the compounds investigated. Thus, several compounds showed a visibly stronger inhibitory effect on the transcription of VEGF mRNA than resveratrol, which lowered *VEGF* gene expression to 65% of the control value. The reduction degrees were particularly strong for compounds **3b** and **17b**, which caused inhibition of the gene expression to 24% and 23%, respectively, of the control value.

The results displayed in Figs. 4 and 5 do not show correlation between the VEGF amount excreted to the medium and the degree of gene expression. Compounds **3b** and **17b**, for instance, proved highly effective in the inhibition of the *VEGF* gene expression but much less so in decreasing the amount of VEGF secreted to the medium. This suggests that these compounds exert the control of VEGF production at a phase different from that of gene transcription, perhaps during the post-translational stage, during which the lighter isoforms such as VEGFA-165 are formed and secreted to the medium [30].

As in the previous case, a separate study was performed for compound **21b**. The inhibition of the *VEGF* gene expression was measured in this case on both HT-29 and MCF-7 cancer cell lines, the results being depicted in Figure 6. When HT-29 cells were incubated in the presence of 1, 2 or 3 ng/mL of compound **21b**, the level of *VEGF* gene expression was similar to that achieved with resveratrol at a

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concentration of 10 μg/mL In other words, the potency of compound **21b** in downregulating *VEGF* gene expression is three orders of magnitude higher than that of resveratrol. In addition, when MCF-7 cells were treated with 2 ng/mL of compound **21b**, only 36 % of VEGF gene expression was detected. Thus, compound **21b** is about 5000 times more active than resveratrol.
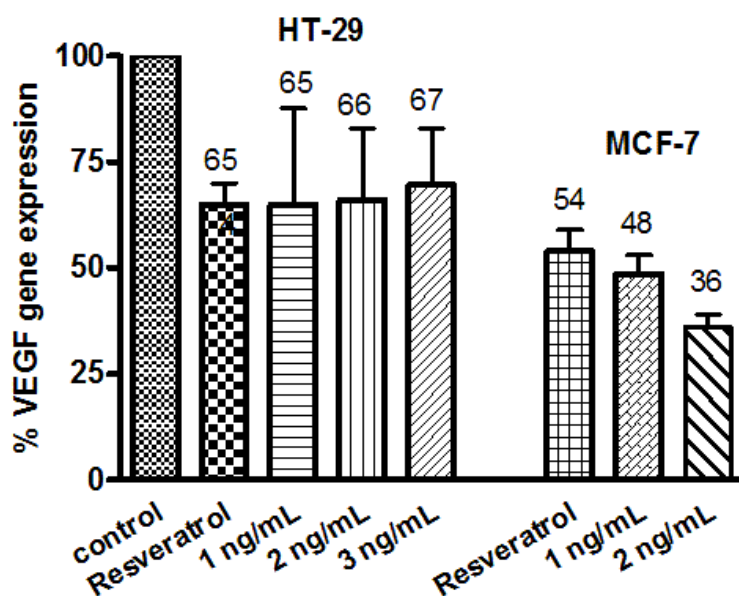
Figure 6. Expression percentage of the *VEGF* gene after 48 h of incubation of HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 μg/mL. Error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

4.3. Effect of stilbene derivatives on the inhibition of the hTERT and c-Myc genes

Human telomerase contains an RNA component (hTERC) that serves as a template for the addition of the repeat nucleotide sequences and also a reverse transcriptase subunit (hTERT) which catalyzes the nucleotide polymerization process. In addition, there are other associated protein factors, the role of which has not yet been completely elucidated. Human telomerase is regulated during development and differentiation, mainly through transcriptional control of the *hTERT* gene, the expression of which is restricted to cells that exhibit telomerase activity. This indicates that the hTERT subunit is the rate limiting factor of the enzyme complex. For the expression of the *hTERT* gene, several transcriptional factors such as c-Myc, a proto-oncoprotein [32], have been found to play an important role through

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upregulation of the mRNA encoding the hTERT protein subunit [33]. Thus, and as an initial study of the potential activity of these stilbene derivatives as inhibitors of telomerase activation, we have investigated their ability to inhibit the expression of the *hTERT* and *c-Myc* genes, which are responsible for the production of the corresponding proteins.

In order to determine whether the studied compounds were able to regulate the expression of the two aforementioned genes, we have performed a RT-qPCR analysis using again HT-29 tumoral cells. The cells were incubated for 48 h in the presence of DMSO (control) and 10 µg/mL of each of the studied compounds (lower than the IC_{50} values). Results are depicted in Fig. 7 (*hTERT* gene expression) and Fig. 8 (*c-Myc* gene expression). As in previous experiments, the highly cytotoxic compound **21b** was the subject of a separate study, which will be described below.

Figure 7. Expression percentage of the *hTERT* gene after 48 h of incubation with HT-29 cells as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

Figure 8. Expression percentage of the *c-Myc* gene after 48 h of incubation with HT-29 cells as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 μg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

As shown in Fig. 7, compounds **10a-12a, 3b, 7b, 14b, 16b** and **22b** showed the highest ability to downregulate the expression of the *hTERT* gene, their activities being somewhat higher than that of resveratrol. As regards the expression of the *c-Myc* gene (Fig. 8), more or less the same compounds showed the highest inhibitory activity. In any case, however, differences in activity with resveratrol were perceptible but not particularly worth noting.

Inhibition of *h-TERT* and *c-Myc* gene expression for compound **21b** was measured on both HT-29 and MCF-7 cancer cell lines. Results are shown in Figs. 9 and 10, which show that compound **21b** was able to downregulate the expression of the *h-TERT* and *c-Myc* genes. The expression of the *h-TERT* gene was decreased to 48% of the control value when 3 ng/mL concentration was used on HT-29 cells, and to 51% when 2 ng/mL concentration was used on MCF-7 cells. Again, the potency of compound **21b** in downregulating the *h-TERT* and *c-Myc* genes is 3000-5000 times greater than that of resveratrol.

Figure 9. Expression percentage of the *hTERT* gene after 48 h of incubation with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

Figure 10. Expression percentage of the *c-Myc* gene after 48 h of incubation of HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent

mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

Table 3 summarizes the results observed for the synthetic compounds **12a**, **3b**, **7b**, **14b**, **16b**, **17b**, **21b** and **22b**, as well as for resveratrol. As regards inhibition of the expression of the *VEGF*, *hTERT* and *c-Myc* genes, stilbene derivatives bearing amino groups (*b*-series) feature much better than those lacking them (*a*-series). Within stilbenes carrying an amide function, the most active is compound **22b** whereas within amino stilbenes, those having a methoxy group (**14b**, **16b**, **17b** and **21b**) perform better than those having a hydroxy group. A very particular case is compound **21b**, which is able to downregulate the secretion of VEGF protein and the expression of *VEGF*, *h-TERT* and *c-Myc* genes at concentrations in the low nanomolar level.

Table 3. Compared percentages of excretion of the VEGF protein and of the expression of the *VEGF*, *hTERT* and *c-Myc* genes.[a]

Compd	Cytotoxicity (IC_{50}, µg/mL)	% Secretion of VEGF protein	% Expression of *VEGF* gene	% Expression of *hTERT* gene	% Expression of *c-Myc* gene
Resv.	34.1 ± 1.4	78	65	57	51
12a	22 ± 4	73	38	48	46
3b	4.4 ± 1.1	78	24	65	46
7b	30.4 ± 1.8	76	38	52	37
14b	21.2 ± 1.5	51	100	35	79
16b	25.3 ± 1	75	67	52	45
17b	19.6 ± 2	73	23	100	34
21b[b]	0.0036 ± 0.001	59	67	48	69
22b	2.5 ± 0.7	79	37	53	45

[a]All values refer to HT-29 cells. The values observed for the control (DMSO without added drugs) are taken as the 100% reference of protein excretion or gene expression. Therefore, the lower the value, the higher the activity of the drug in the corresponding activity. Except for **21b**, measurements were performed at concentration values of 10 µg/mL.
[b]At a concentration of 3 ng/mL.

The biological activities of resveratrol are often ascribed, at least in part, to its antioxidant activity. The various biological activities of resveratrol analogues depend upon the number and location of the

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hydroxy groups [34], minor structural changes in such stilbenes having major effects on their biological activity [35]. In the present study, a series of 39 stilbenes carrying hydroxy, methoxy and amino groups were compared with resveratrol for their cytotoxicity and for their ability to inhibit VEGF secretion and the expression of the *VEGF*, *h-TERT* and *c-Myc* genes. Several compounds were found to be more active than resveratrol, the case of (*E*)-4-(4-methoxystyryl)aniline (**21b**) being particularly interesting because of its high cytotoxicity. In order to find some relationships between chemical structure and biological activity, we have graphically represented α coefficients vs. β coefficients of the studied compounds (Fig. 11). Figs. 11a to 11c display the α and β values for stilbenes carrying an *ortho-*, *meta-* and *para*-hydroxy group, respectively. Figure 11d shows values for the remaining compounds.
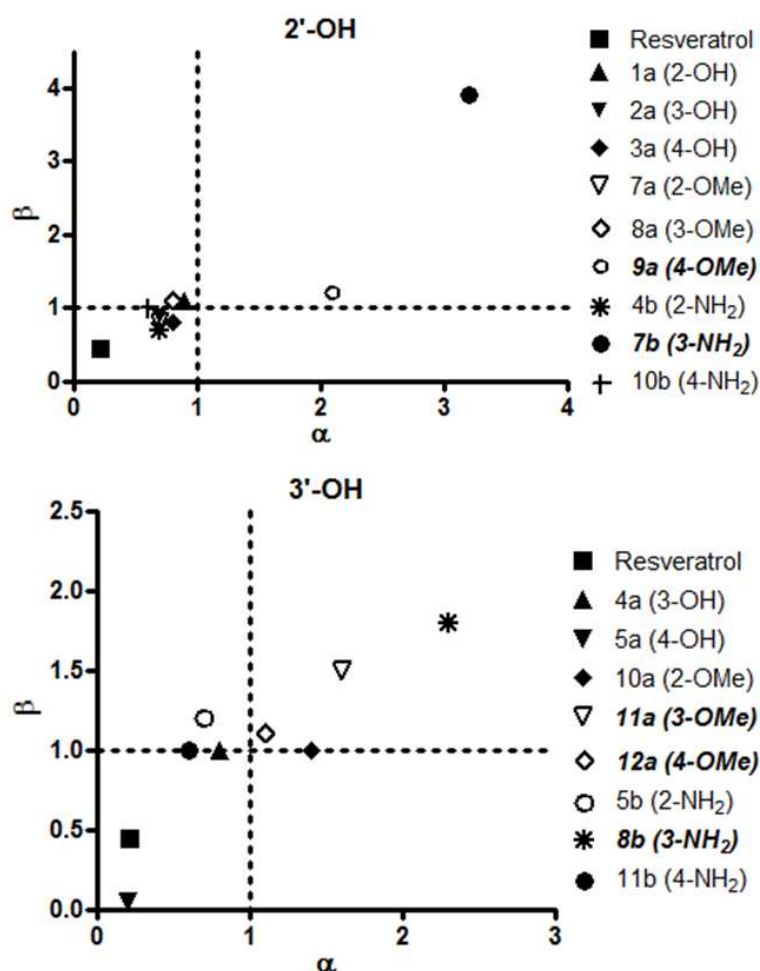
Figure 11. α coefficients vs. β coefficients of the synthetic stilbenes.
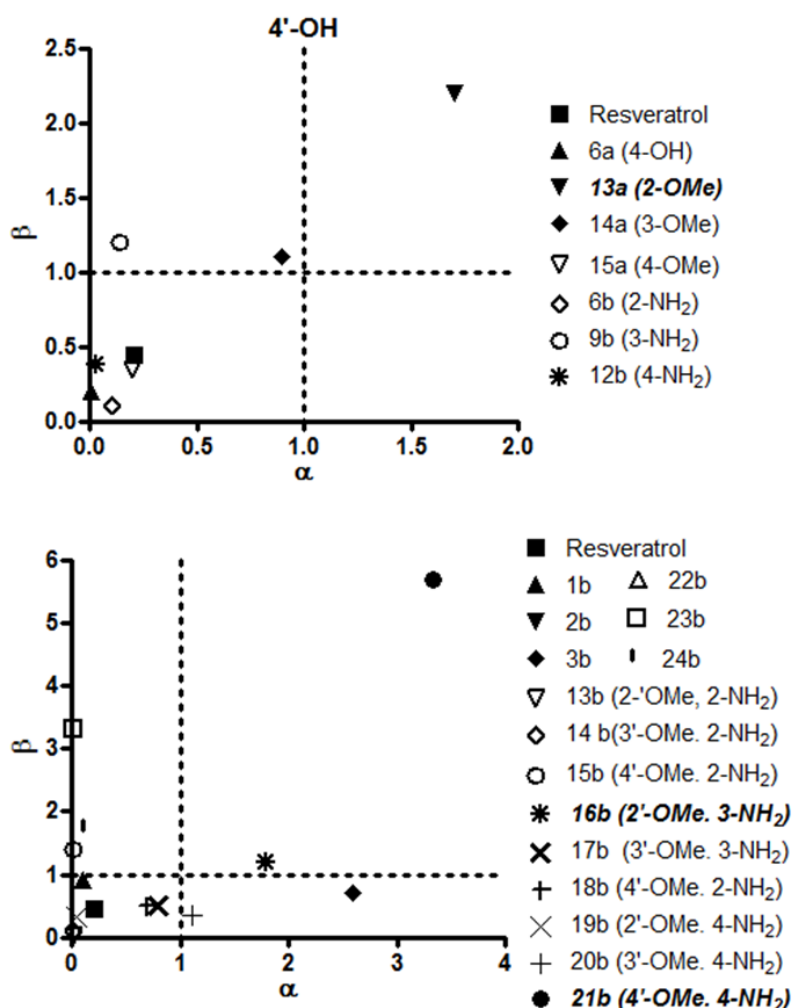
Figure 11 (cont.)**.** α coefficients vs. β coefficients of the synthetic stilbenes.

As mentioned above, the higher the coefficient values, the better the safety margin for the corresponding compound from the therapeutic point of view. Aside from compound **21b**, which is three orders of magnitude more cytotoxic than the others, the best safety margins are achieved with compounds **9a** (2'-OH, 4-OMe), **13a** (4'-OH, 2-OMe), **7b** (2'-OH, 3-NH_2), **8b** (3'-OH, 3-NH_2) and **16b** (2'-OMe, 3-NH_2). These three amino stilbenes share in common an amino group in *meta* position, regardless of the position occupied by the hydroxy or methoxy group. Compound **21b**, however, which is cytotoxic at nanomolar level and shows the highest safety margins, has its amino group in the *para* position. This may possibly indicate that, in comparison with the rest of compounds, **21b** shows a different mode of action.

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In Figure 12, the percentage of VEFG protein secretion is graphically represented vs. the percentage of *h-TERT* gene expression. Compounds **14b** and **21b** are located in the lower left quadrant, where high inhibition of VEGF secretion and high inhibition of *h-TERT* gene expression are combined. It is worth noting that this desirable dual mode of action is exhibited by compound **21b** at low nanomolar concentrations, whereas compound **14b** achieves it only at the micromolar level.
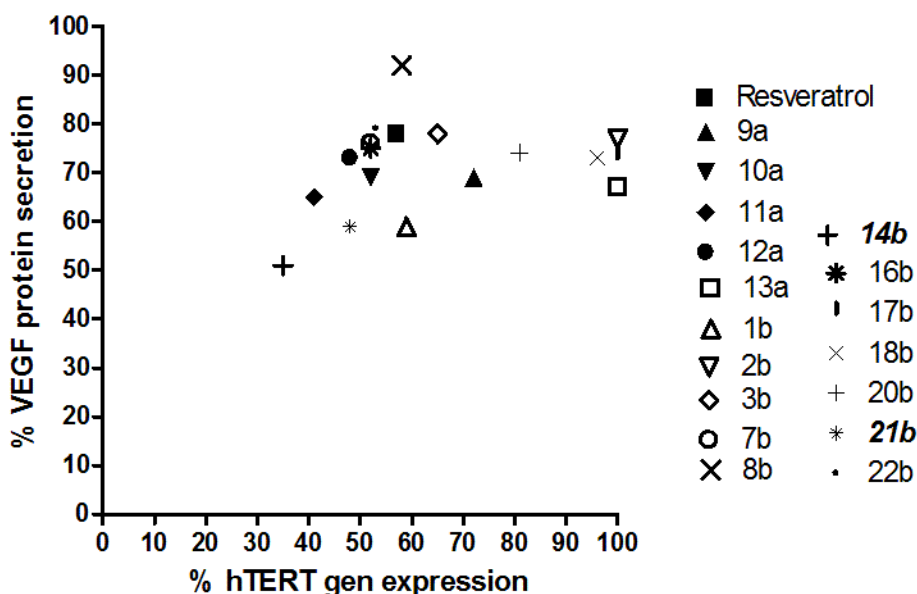
Figure 12. Percentage of VEFG protein secretion vs. percentage of *h-TERT* gene expression.

5. Summary and conclusions

In conclusion, 39 stilbene derivatives have been synthesized and evaluated for their biological activity. One of them, (*E*)-4-(4-methoxystyryl)aniline (compound **21b**) has proven particularly interesting. Not only has this compound a strong cytotoxicity at the low nanomolar level and a good safety margin, it further combines these features with an ability to inhibit VEFG protein secretion and the expression of the telomerase related *h-TERT* and *c-Myc* genes at similarly low concentrations. Compound **21b** therefore may possibly display anticancer activity through a multiple mode of action. It is worth noting that **21b** shows an extremely high difference in activity with all other compounds, including its positional isomers **19b** and **20b**, and its demethyl derivative **12b**. This may possibly suggest that its interactions with its (still unknown) biomolecular targets, and perhaps also the targets themselves, are of a different nature. More studies will be required to acquire a knowledge about its mechanism of action.

6. Experimental

6.1. Chemistry. General procedures

The general reaction conditions and the physical and spectral data of all synthetic intermediates and final compounds are described in detail in the Supporting Information. The samples of compounds used for the biological studies were purified to > 95% by means of preparative HPLC.

6.2. Biological studies. Materials and methods

6.2.1. Reagents and cell culture

Cell culture media were purchased from Gibco (Grand Island, NY, USA). Fetal bovine serum (FBS) was a product of Harlan-Seralab (Belton, U.K.). Supplements and other chemicals not listed in this section were obtained from Sigma Chemicals Co. (St. Louis, Mo., USA). Plastics for cell culture were supplied by Thermo Scientific™ BioLite. All tested compounds were dissolved in DMSO at a concentration of 10 µg/mL and stored at –20°C until use.

Cell lines were maintained in Dulbecco's modified Eagle's medium (DMEM) containing glucose (1 g/L), glutamine (2 mM), penicillin (50 IU/mL), streptomycin (50 µg/mL) and amphoterycin (1.25 µg/mL), supplemented with 10% FBS.

6.2.2. Cytotoxicity assays

The 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; Sigma Chemical Co., St. Louis, MO) dye reduction assay in 96-well microplates was used, as previously described [36]. Some 5 x 10^3 cells of HT-29, MCF-7 or HEK-293 cells in a total volume of 100 µL of their respective growth media were incubated with serial dilutions of the tested compounds. After 3 days of incubation (37 °C, 5% CO_2 in a humid atmosphere), 10 µl of MTT (5 mg/ml in PBS) were added to each well and the plate was incubated for further 4 h (37 °C). The resulting formazan was dissolved in 150 µL of 0.04 N HCl/2-propanol and read at 550 nm. All determinations were carried out in triplicate.

6.2.3. ELISA analysis

HT-29 cells at 70–80% confluence were collected after serum starvation for 24 h. Cells were incubated with the selected stilbene derivatives in DMSO for 72 h (Fig. 3). Culture supernatants were collected and VEGF secreted by HT-29 cells was determined using Invitrogen Human Vascular Endothelial Growth Factor ELISA Kit according to the manufacturer's instructions.

6.2.4. RT-qPCR analysis

HT-29 cells at 70–80% confluence were collected after serum starvation for 24 h. Cells were incubated with 10 µg/mL of the corresponding drugs in DMSO for 48 h. Cells were collected and the total cellular RNA from HT-29 cells was isolated using Ambion RNA extraction Kit according to the manufacturer's instructions. The cDNA was synthesized by MMLV-RT with 1-21 µg of extracted RNA and oligo(dT)15 according to the manufacturer's instructions.

Amplification of the genes was performed by use of a StepOnePlus™ thermalcycler. Fast TaqMan Gene Expression Master Mix containing the appropriate buffer for the amplification conditions, dNTPs, thermostable DNA polymerase enzyme and a passive reference probe was used. Each of the genes were amplified using predesigned primers by Life Technologies TaqMan® Gene Expression Assays, Hs99999903-m1 (β-actin), Hs00900055-m1 (VEGF), Hs00972646-m1 (*hTERT*) and Hs00153408-m1 (*c-Myc*).

6.2.5. Statistical analysis

Data are expressed as the mean ± SEM. Statistical analyses were done using Microsoft Excel and GraphPad Prism®. Differences between means were determined using Student's t-Test or one-way ANOVA with Dunnett's Multiple Comparison Test, and considered to be statistically significant at p≤0.001.

Supplementary Information

Description of general features and reaction conditions. Spectral data and graphical NMR spectra of all new compounds.

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List of captions

Figure 1. Structure of resveratrol

Figure 2. Structures of synthetic resveratrol analogues.

Scheme 1. Synthesis of stilbene derivatives by means of palladium-catalyzed Heck couplings (for more detailed descriptions of the reaction conditions and yields, see Supporting Information).

Table 1. IC_{50} values (µg/mL) and selectivity coefficients for compounds of the *a*-series.

Table 2. IC_{50} values (µg/mL) and selectivity coefficients for compounds of the *b*-series.

Figure 3. VEGF protein secretion from HT-29 cells determined by means of the ELISA procedure. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 4. VEGF protein secretion from HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 5. Expression percentage of the *VEGF* gene after 48 h of incubation of HT-29 cells determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 6. Expression percentage of the *VEGF* gene after 48 h of incubation of HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 7. Expression percentage of the *hTERT* gene after 48 h of incubation with HT-29 cells as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 8. Expression percentage of the *c-Myc* gene after 48 h of incubation with HT-29 cells as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of all compounds was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests ($P < 0.001$).

Figure 9. Expression percentage of the *hTERT* gene after 48 h of incubation with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

Figure 10. Expression percentage of the *c-Myc* gene after 48 h of incubation of HT-29 and MCF-7 cells with compound **21b** as determined by means of the RT-qPCR methodology. At least three measurements were performed in each case. Concentration of resveratrol was 10 µg/mL. Bars represent mean values of VEGF secretion (percentage values related to control) and error bars indicate standard errors of the mean. The statistical significance was evaluated using one-sample t-tests (P < 0.001).

Table 3. Compared percentages of excretion of the VEGF protein and of the expression of the *VEGF*, *hTERT* and *c-Myc* genes.[a]

Figure 11. α coefficients vs. β coefficients of the synthetic stilbenes.

Figure 11 (cont.)**.** α coefficients vs. β coefficients of the synthetic stilbenes.

Figure 12. Percentage of VEFG protein secretion vs. percentage of *h-TERT* gene expression.
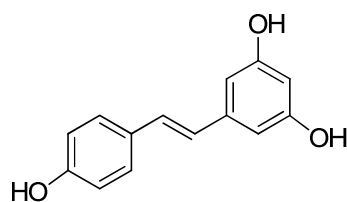
Figure 1. Structure of resveratrol

1a	R^1 = 2-OH	R^2 = 2´-OH	**6a**	R^1 = 4-OH	R^2 = 4´-OH	**11a** R^1 = 3-OMe R^2 = 3´-OH
2a	R^1 = 3-OH	R^2 = 2´-OH	**7a**	R^1 = 2-OMe	R^2 = 2´-OH	**12a** R^1 = 4-OMe R^2 = 3´-OH
3a	R^1 = 4-OH	R^2 = 2´-OH	**8a**	R^1 = 3-OMe	R^2 = 2´-OH	**13a** R^1 = 2-OMe R^2 = 4´-OH
4a	R^1 = 3-OH	R^2 = 3´-OH	**9a**	R^1 = 4-OMe	R^2 = 2´-OH	**14a** R^1 = 3-OMe R^2 = 4´-OH
5a	R^1 = 4-OH	R^2 = 3´-OH	**10a**	R^1 = 2-OMe	R^2 = 3´-OH	**15a** R^1 = 4-OMe R^2 = 4´-OH

1b	R^1 = 2-NH_2	R^2 = H	**9b**	R^1 = 3-NH_2	R^2 = 4´-OH	**17b** R^1 = 3-NH_2 R^2 = 3´-OMe
2b	R^1 = 3-NH_2	R^2 = H	**10b**	R^1 = 4-NH_2	R^2 = 2´-OH	**18b** R^1 = 3-NH_2 R^2 = 4´-OMe
3b	R^1 = 4-NH_2	R^2 = H	**11b**	R^1 = 4-NH_2	R^2 = 3´-OH	**19b** R^1 = 4-NH_2 R^2 = 2´-OMe
4b	R^1 = 2-NH_2	R^2 = 2´-OH	**12b**	R^1 = 4-NH_2	R^2 = 4´-OH	**20b** R^1 = 4-NH_2 R^2 = 3´-OMe
5b	R^1 = 2-NH_2	R^2 = 3´-OH	**13b**	R^1 = 2-NH_2	R^2 = 2´-OMe	**21b** R^1 = 4-NH_2 R^2 = 4´-OMe
6b	R^1 = 2-NH_2	R^2 = 4´-OH	**14b**	R^1 = 2-NH_2	R^2 = 3´-OMe	**22b** R^1 = 2-$NHCO(CH_2)_{10}Me$ R^2 = H
7b	R^1 = 3-NH_2	R^2 = 2´-OH	**15b**	R^1 = 2-NH_2	R^2 = 4´-OMe	**23b** R^1 = 3-$NHCO(CH_2)_{10}Me$ R^2 = H
8b	R^1 = 3-NH_2	R^2 = 3´-OH	**16b**	R^1 = 3-NH_2	R^2 = 2´-OMe	**24b** R^1 = 4-$NHCO(CH_2)_{10}Me$ R^2 = H

Figure 2. Structures of synthetic resveratrol analogues.

Scheme 1. Synthesis of stilbene derivatives by means of palladium-catalyzed Heck couplings (for more detailed descriptions of the reaction conditions and yields, see Supporting Information).

Table 1. IC$_{50}$ values (µg/mL) and selectivity coefficients for compounds of the *a*-series.

Compound	HT-29	MCF-7	HEK-293	α[b]	β[c]
Resveratrol	34.1 ± 1.4	16.1 ± 1.1	7.1 ± 1.0	0.2	0.4
1a	25 ± 7	23.0 ± 1.3	24.3 ± 2.3	0.9	1.1
2a	27 ± 7	20.1 ± 2.1	18 ± 8	0.7	0.9
3a	22 ± 9	21 ± 3	17.5 ± 1.5	0.8	0.8
4a	25.9 ± 1.0	21 ± 6	21 ± 5	0.8	1
5a	6.9 ± 1.9	23 ± 3	1.1 ± 0.7	0.2	< 0.1
6a	>100	6.8 ± 0.2	1.2 ± 0.1	< 0.1	0.2
7a	23 ± 6	17.9 ± 2.4	15.0 ± 2.5	0.7	0.8
8a	26 ± 3	18 ± 4	20 ± 4	0.8	1.1
9a	10 ± 4	17 ± 6	21 ± 4	2.1	1.2
10a	17 ± 4	24.3 ± 1.0	24 ± 3	1.4	1
11a	11.5 ± 3	13 ± 3	19 ± 7	1.6	1.5
12a	22 ± 4	21 ± 5	24 ± 3	1.1	1.1
13a	24 ± 3	19 ± 4	41 ± 3	1.7	2.2
14a	26 ± 4	21.0 ± 1.5	24 ± 3	0.9	1.1
15a	17.4 ± 0.2	11.3 ± 2.3	3.9 ± 0.5	0.2	0.3

[a]Values are the average (± s.d.) of three different measurements performed as described in the Material and Methods section. [b]α = IC$_{50}$ (HEK-293) / IC$_{50}$ (HT-29). [c]β = IC$_{50}$ (HEK-293) / IC$_{50}$ (MCF-7). Values of α and β have been rounded off to a decimal figure.

Table 2. IC_{50} values (µg/mL) and selectivity coefficients for compounds of the *b*-series.

Compound	HT-29	MCF-7	HEK-293	α^b	β^c
Resveratrol	34.1 ± 1.4	16.1 ± 1.1	7.1 ± 1.0	0.2	0.4
1b	17 ± 4	1.4 ± 0.1	1.3 ± 0.2	0.1	0.9
2b	12.9 ± 0.9	14 ± 4	84 ± 11	6.5	6
3b	4.4 ± 1.1	16 ± 0.1	11.4 ± 0.4	2.6	0.7
4b	36 ± 5	39 ± 3	27 ± 7	0.7	0.7
5b	30 ± 5	17 ± 3	20 ± 7	0.7	1.2
6b	>100	79 ± 3	7 ± 3	< 0.1	< 0.1
7b	30.4 ± 1.8	24.7 ± 0.2	96 ± 15	3.2	3.9
8b	43 ± 16	54 ± 7	>100	2.3	1.8
9b	51 ± 16	6 ± 3	7 ± 3	0.1	1.2
10b	37 ± 5	23 ± 4	21.4 ± 1.4	0.6	1
11b	42 ± 6	24.8 ± 0.4	25 ± 8	0.6	1
12b	30.6 ± 2.4	2.2 ± 0.4	0.8 ± 0.1	0.03	0.4
13b	>100	>100	2.1 ± 0.7	< 0.1	< 0.1
14b	21.2 ± 1.5	6.1 ± 1.8	0.5 ± 0.1	< 0.1	< 0.1
15b	73 ± 5	0.4 ± 0.1	0.6 ± 0.1	< 0.1	1.4
16b	25.3 ± 0.1	38.7 ± 0.6	45 ± 0.3	1.8	1.2
17b	19.6 ± 2.0	34 ± 5	16 ± 5	0.8	0.5
18b	16.3 ± 0.4	23 ± 4	11 ± 0.3	0.7	0.5
19b	22 ± 4	3.2 ± 1.7	1 ± 0.3	< 0.1	0.3
20b	16.1 ± 0.1	52 ± 6	18 ± 4	1.1	0.3
21b	0.0036 ± 0.0015	0.0021 ± 0.0005	0.012 ± 0.004	3.4	5.7
22b	2.5 ± 0.7	12 ± 0.9	16 ± 9	6.4	1.3
23b	>100	3.5 ±0.5	11.6 ± 0.6	< 0.1	3.3
24b	>100	5.8 ± 0.1	10 ± 4	< 0.1	1.7

[a]Values are the average (± s.d.) of three different measurements performed as described in the Material and Methods section. [b]α = IC_{50} (HEK-293) / IC_{50} (HT-29). [c]β = IC_{50} (HEK-293) / IC_{50} (MCF-7). Values of α and β have been rounded off to a decimal figure.

Figure 3.

Figure 4.

Figure 5.
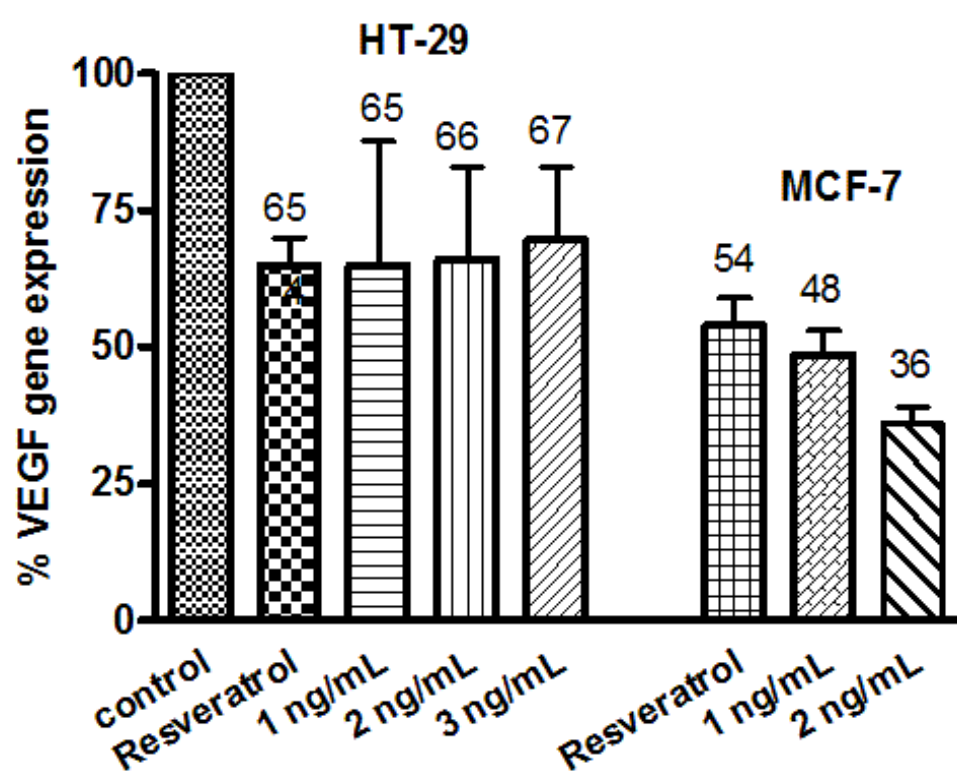
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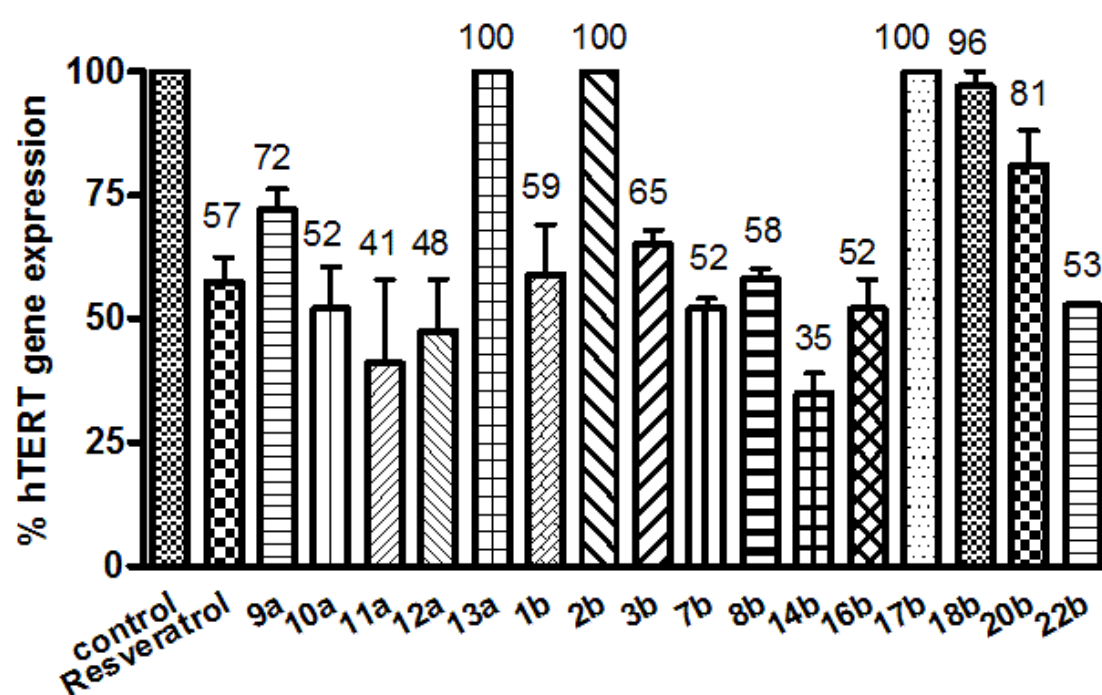
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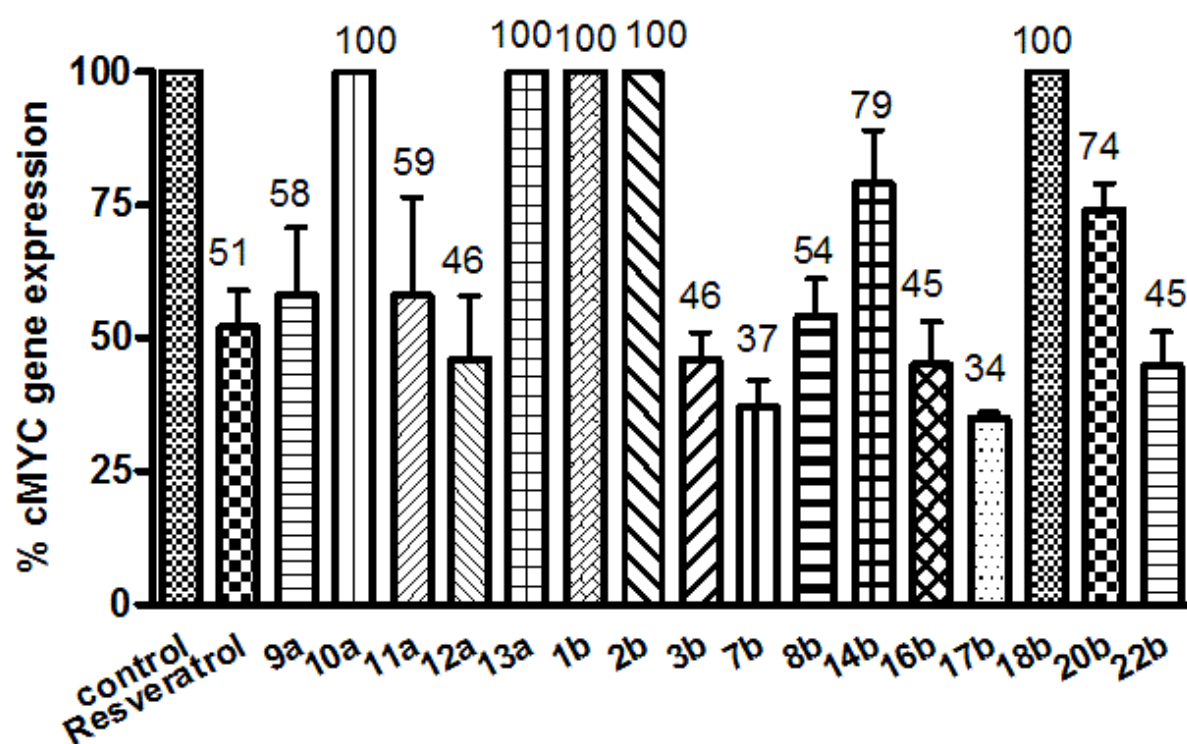
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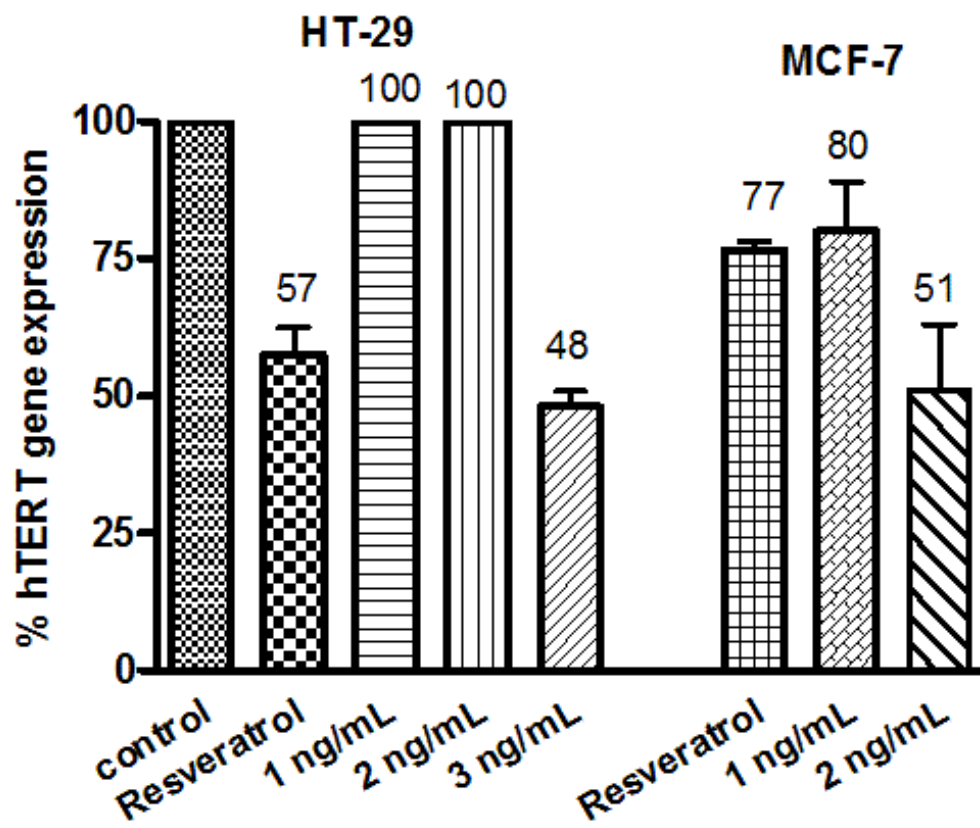
Figure 9.
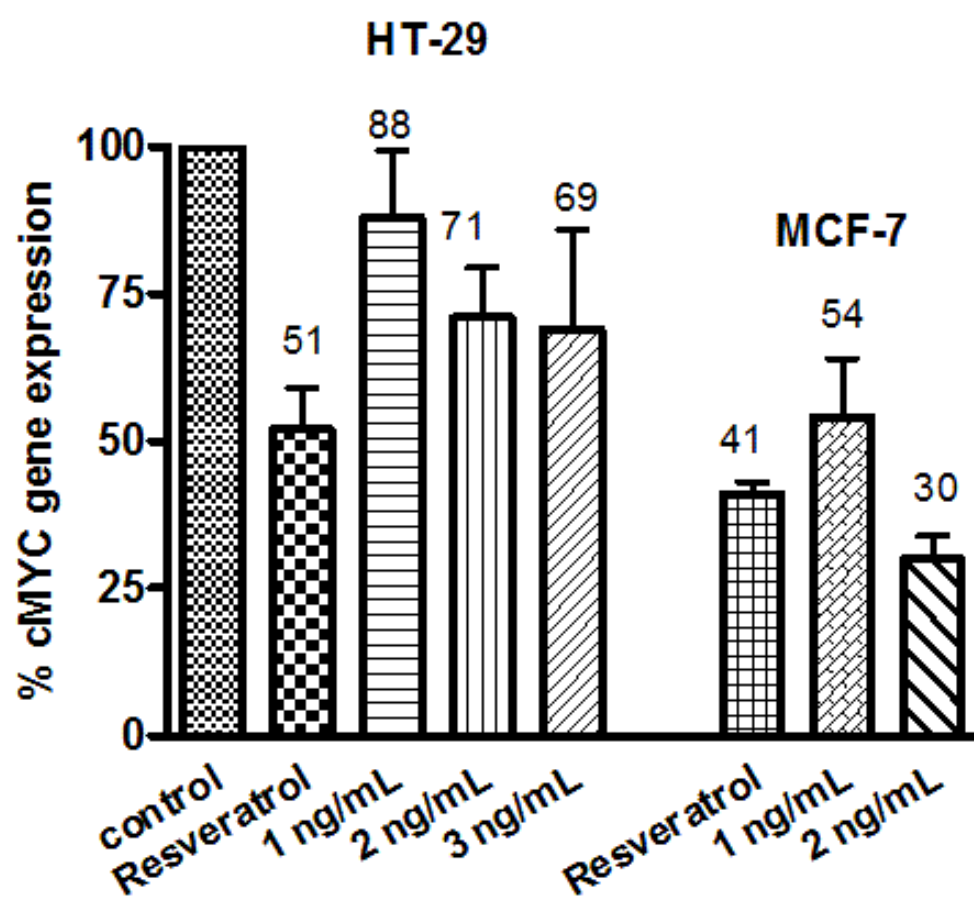
Figure 10.

Table 3. Compared percentages of excretion of the VEGF protein and of the expression of the *VEGF*, *hTERT* and *c-Myc* genes.[a]

Compd	Cytotoxicity (IC$_{50}$, µg/mL)	% Secretion of VEGF protein	% Expression of *VEGF* gene	% Expression of *hTERT* gene	% Expression of *c-Myc* gene
Resv.	34.1 ± 1.4	78	65	57	51
12a	22 ± 4	73	38	48	46
3b	4.4 ± 1.1	78	24	65	46
7b	30.4 ± 1.8	76	38	52	37
14b	21.2 ± 1.5	51	100	35	79
16b	25.3 ± 1	75	67	52	45
17b	19.6 ± 2	73	23	100	34
21b[b]	0.0036 ± 0.001	59	67	48	69
22b	2.5 ± 0.7	79	37	53	45

[a]All values refer to HT-29 cells. The values observed for the control (DMSO without added drugs) are taken as the 100% reference of protein excretion or gene expression. Therefore, the lower the value, the higher the activity of the drug in the corresponding activity. Except for **21b**, measurements were performed at concentration values of 10 µg/mL.
[b]At a concentration of 3 ng/mL.

Figure 11.

45
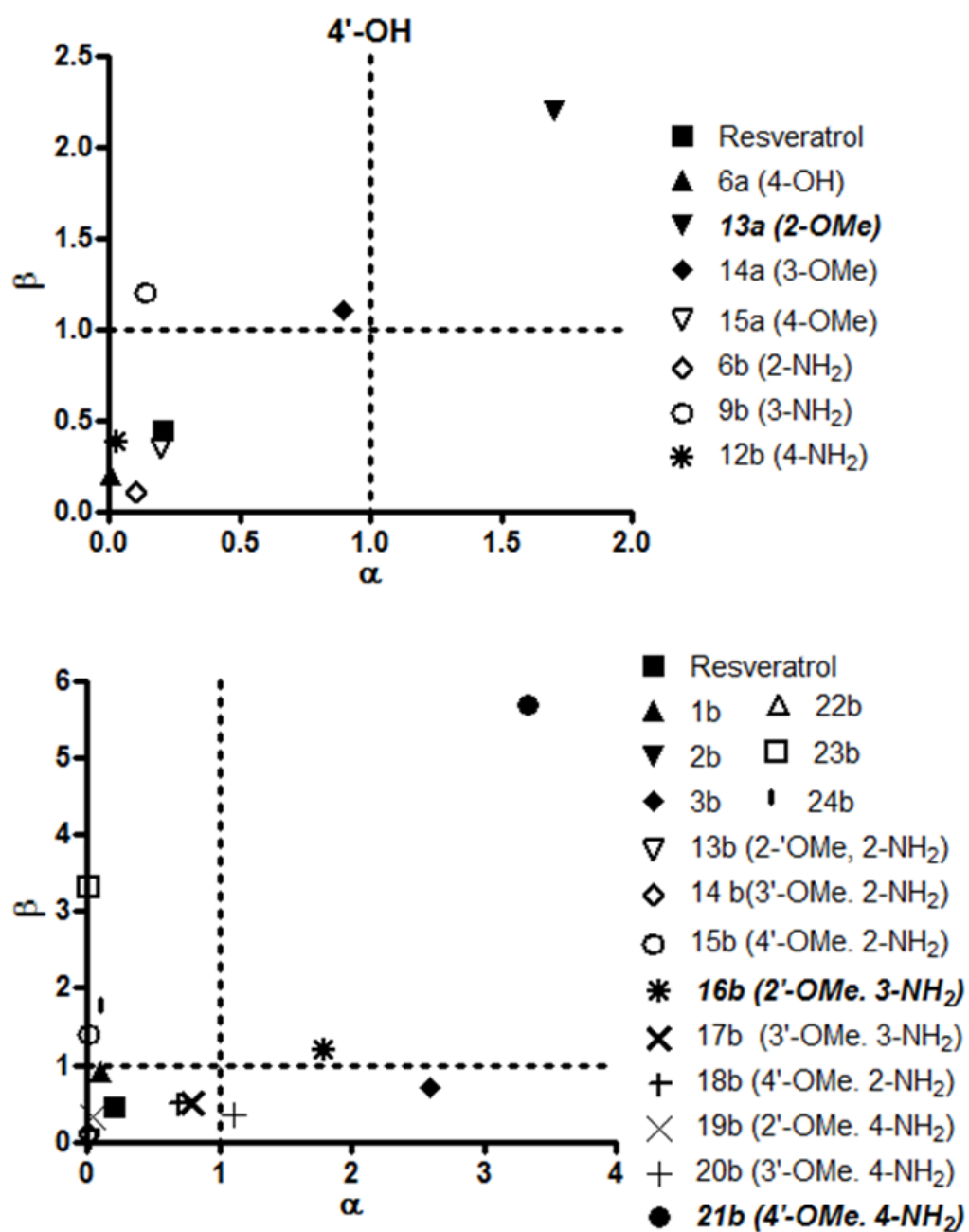
Figure 11 (cont.).
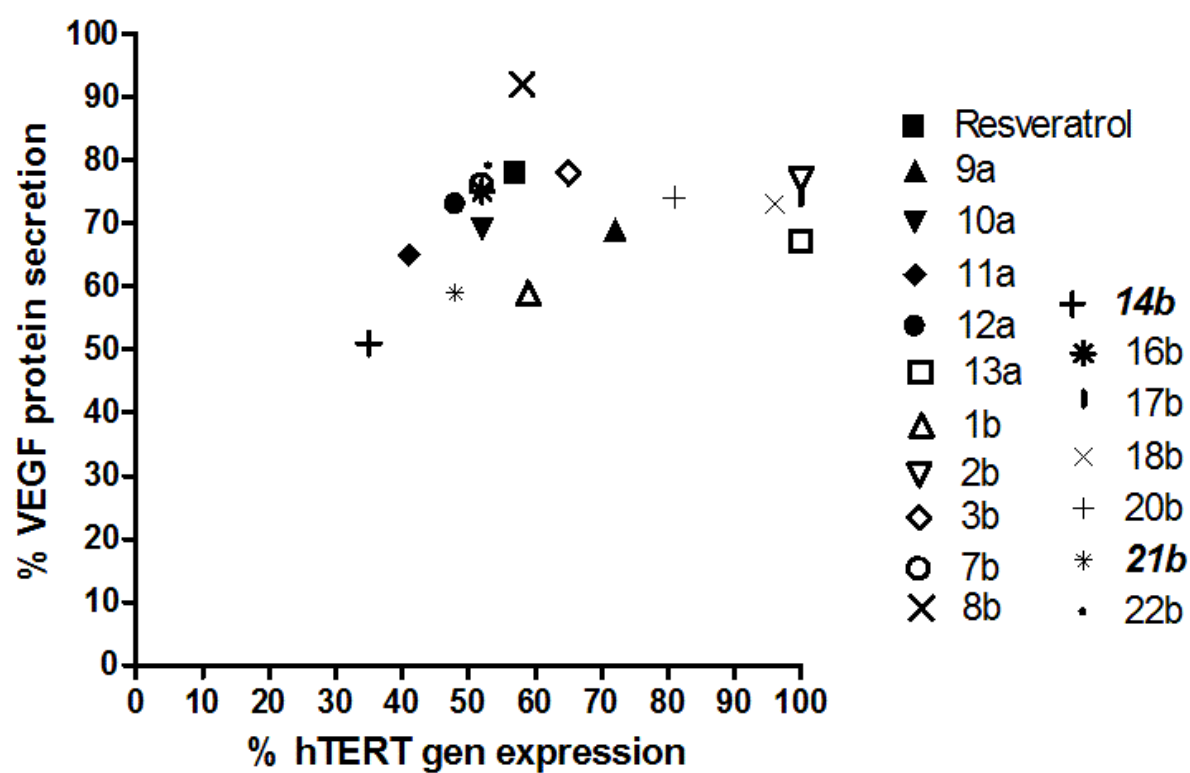
Figure 12.

Inhibitory effect of cytotoxic stilbenes related to resveratrol on the expression of the *VEGF*, *hTERT* and *c-Myc* genes

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Supporting Information

Contents:

General features. NMR spectra were measured at 25 °C. The signals of the deuterated solvent (CDCl$_3$, CD$_3$OD or DMSO-d$_6$) were taken as the reference. Multiplicity assignments of ^{13}C signals were made by means of the DEPT pulse sequence. High resolution mass spectra were run by the electrospray mode (ESMS). In some cases where no positive ions were produced, negative ions were measured instead. IR data were measured with oily films on NaCl plates (oils) or KBr pellets (solids) and are given only for molecules with relevant functional groups (OH, C=O, NH$_2$). Melting points are not corrected. Experiments which required an inert atmosphere were carried out under dry N$_2$ in a flame-dried glassware. THF was freshly distilled from sodium/benzophenone ketyl and transferred via syringe. Dimethylformamide was freshly distilled from CaH$_2$. Commercially available reagents were used as received. Where solutions were filtered through a Celite pad, the pad was additionally washed with the same solvent used, and the washings incorporated to the main organic layer. Acronyms used hereafter: DMF = *N,N*-dimethylformamide, TBAB = tetra-*n*-butylammonium bromide.

General reaction conditions of Heck couplings
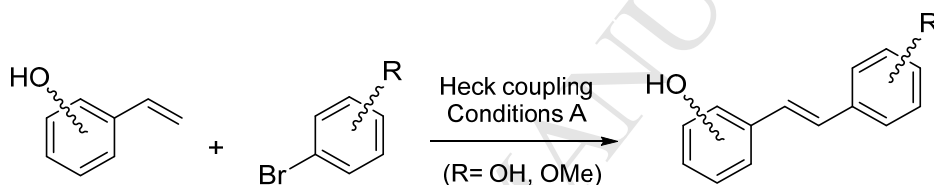
Heck coupling - Conditions A[1]

The appropriate bromoderivative (1 mmol) was placed in a 10-mL glass tube together with the corresponding styrene derivative (1.5 mmol), piperidine (0.36 mL, 3.65 mmol), Pd(PPh$_3$)$_4$ (0.035 g, 0.03 mmol) and DMF (2 mL). In some cases, addition of LiCl (0.0034 g, 0.08 mmol) was found necessary. The vessel was sealed with a septum, shaken, and placed into the microwave cavity. Microwave irradiation was initiated with a potency of 150 W, the temperature being then increased from room temperature to 180 ºC. Once this value was reached, the reaction mixture was held at this temperature for 3 h. After allowing the mixture to cool down to room temperature, the reaction vessel was opened and the mixture was filtered through a Celite pad. The filtrate was subsequently made acid (aqueous HCl, pH 5-6) or else basic (aqueous NaOH, pH 8-9). The mixture was poured into a separation funnel and extracted with ethyl ether (3 x 40 mL). The combined organic layers were washed with aqueous NH$_4$Cl (20 mL), water (20 mL) and brine (20 mL). After desiccation over anhydrous Na$_2$SO$_4$, the volatiles were removed under reduced pressure. This afforded an oily material which was subjected to column chromatography on silica gel (hexane-EtOAc mixtures) to yield the desired stilbene derivative.

Conditions A: Pd(P(Ph$_3$)$_4$), piperidine, DMF, 180 ºC (MW 150 W), 3 h. Yields: **1a,** 61%; **2a,** 62%; **3a**, 67%; **4a,** 81%; 5a, 88%; **6a**, 64%; **7a**, 56%; **8a**, 76%; **9a**, 52%; **10a**, 85%, **11a**, 71%; **12a,** 72%; **13a**, 53%; **14a**, 62%; **15a**, 50%; **4b**, 64%; **5b**, 62%; **7b**, 70%, **9b**, 75%.

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[1] N. Sharma, D. Mohanakrishnan, A. Shard, A. Sharma, Saima, A. K. Sinha, D. Sahal, Stilbene-Chalcone Hybrids: Design, Synthesis, and Evaluation as a New Class of Antimalarial Scaffolds That Trigger Cell Death through Stage Specific Apoptosis , J. Med. Chem. 55 (2012) 297-311.
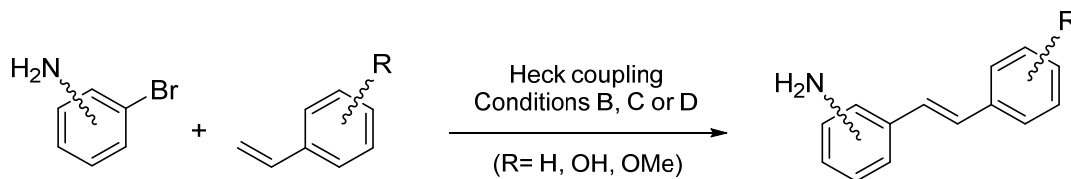
Heck coupling - Conditions B[2]

The appropriate bromo aniline (1 mmol) was placed in a 10-mL glass tube together with 3-vinylphenol (1 mmol), tri-*n*-butylamine (0.22 mL, 1.2 mmol), TBAB (0.387 g, 1.2 mmol) and water (3 mL). The mixture was stirred until homogeneity and then treated with $Pd(NH_3)_2Cl_2$ (2 mg, 0.01 mmol). The vessel was sealed with a septum, shaken, and placed into the microwave cavity. Microwave irradiation was initiated with a potency of 70 W, the temperature being then increased from room temperature to 140 ºC. Once this value was reached, the reaction mixture was held at this temperature for 1 h. After allowing the mixture to cool down to room temperature, the reaction vessel was opened and the mixture was poured into a separation funnel. Water (10 mL) and EtOAc (10 mL) were added and the organic material was extracted. After further extraction of the aqueous layer with EtOAc, the organic layers were combined and dried over anhydrous Na_2SO_4. The volatiles were removed under reduced pressure to afford an oily residue, which was subjected to column chromatography on silica gel (hexane-EtOAc mixtures) to yield the desired stilbene derivative.

Conditions B: $Pd(NH_3)_2Cl_2$, Bu_3N, TBAB, H_2O, 140 ºC (MW 70 W), 1.5 h. Yields: **8b**, 74%; **11b**, 68%.

Heck coupling - Conditions C[1,2]

The appropriate bromo aniline (1 mmol) was placed in a 10-mL glass tube together with the corresponding styrene (1 mmol), piperidine (0.15 mL, 1.5 mmol), $Pd(NH_3)_2Cl_2$ (8.4 mg, 0.04 mmol) and DMF (4 mL). The vessel was sealed with a septum, shaken, and placed into the microwave cavity. Microwave irradiation was initiated with a potency of 70 W, the temperature being then increased from room temperature to 140 ºC. Once this value was reached, the reaction mixture was held at this temperature for 1.5 h. After allowing the mixture to cool down to room temperature, the reaction vessel was opened and the reaction mixture was filtered through a Celite pad with EtOAc. The filtrate was poured into a separating funnel, and washed with ammonium chloride (2 x 10 mL) and then with brine (2 x 10 mL). After desiccation of the organic layer over anhydrous Na_2SO_4, the volatiles were removed under reduced pressure. This afforded an oily material which was subjected to column chromatography on silica gel (hexane-EtOAc mixtures) to yield the desired stilbene derivative.

Conditions C: $Pd(NH_3)_2Cl_2$, piperidine, DMF, 140 ºC (MW 70 W), 1.5 h. Yields: **1b,** 75%; **2b**, 69%; **3b**, 68%; **6b**, 51%; **10b**, 49%; **12b**, 52%; **22b**, 55%; **24b**, 58%.

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[2] S.-H. Huang, J.-R. Chen, F.-Y. Tsai, Palladium(II)/Cationic 2,2 '-Bipyridyl System as a Highly Efficient and Reusable Catalyst for the Mizoroki-Heck Reaction in Water , Molecules 15 (2010) 315-330.

Heck coupling - Conditions D[3]

The appropriate bromo aniline (1 mmol) was placed in a 10-mL glass tube together with the corresponding styrene (1 mmol), $Pd(NH_3)_2Cl_2$ (1.1 mg, 0.005 mmol), K_2CO_3 (0.276 g, 2 mmol) and DMF (1.5 mL). The vessel was sealed with a septum, shaken, and placed into the microwave cavity. Microwave irradiation was initiated with a potency of 25 W, the temperature being then increased from room temperature to 175 ºC. Once this value was reached, the reaction mixture was held at this temperature for 30 min. After allowing the mixture to cool down to room temperature, the reaction vessel was opened and the reaction mixture was filtered through a Celite pad with EtOAc. The filtrate was poured into a separating funnel, and washed with water (2 x 10 mL) and then with brine (2 x 10 mL). After desiccation of the organic layer over anhydrous Na_2SO_4, the volatiles were removed under reduced pressure. This afforded an oily material which was subjected to column chromatography on silica gel (hexane-EtOAc mixtures) to yield the desired stilbene derivative.

Conditions D: $Pd(NH_3)_2Cl_2$, K_2CO_3, DMF, 175 ºC (MW 25 W), 30 min. Yields: **13b**, 72%; **14b**, 69%; **15b**, 59%; **16b**, 71%; **17b**, 73%; **18b**, 71%; **19b**, 74%; **20b**, 45%; **21b**, 55%.

Amide 23b[4]

A stirred solution of dodecanoic acid (200 mg, 1 mmol) and triethylamine (0.15 mL, 1.05 mmol) in dry dichloromethane (5 mL) was treated at 0º C with a solution of chlorosulfonyl isocyanate (0.09 mL, 1 mmol) in dry dichloromethane (5 mL). The reaction mixture was stirred at 0º C for 3 h, followed by addition of **2b** (2 mmol). The reaction mixture was stirred at room temperature for 16 h and then poured into water (20 mL). After extraction of the reaction mixture with EtOAc (3 x 25 mL), the combined organic layers were washed with aqueous sodium hydrogen carbonate (2 x 20 mL) and then with water (2 x 20 mL). Dessication of the organic layer over anhydrous Na_2SO_4 and removal of volatiles under reduced pressure gave an oily residue which was subjected to column chromatography on silica gel (hexane-EtOAc, 9:1) to yield **23b** (64%).

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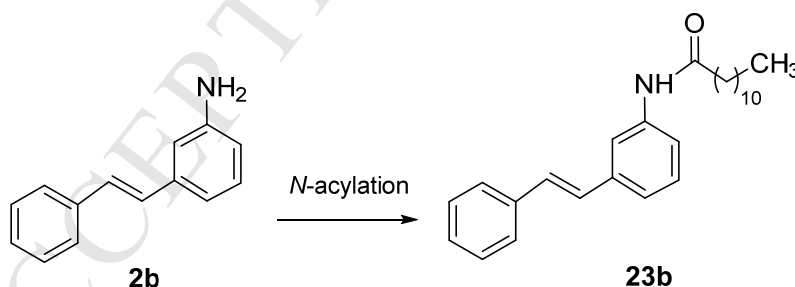
[3] C. M. Kormos, N. E. Leadbeater, Preparation of nonsymmetrically substituted stilbenes in a one-pot two-step Heck strategy using ethene as a reagent , J. Org. Chem. 73 (2008) 3854-3858.

[4] K. S. Keshavamurthy, Y. D. Vankar, D. N. Dhar, Preparation of acid anhydrides, amides, and esters using chlorosulfonyl isocyanate as a dehydrating agent , Synthesis (1982) 506-508.

Analytical data of stilbene derivatives

For the biological assays, samples of synthetic resveratrol analogues have been carefully purified by means of preparative HPLC.
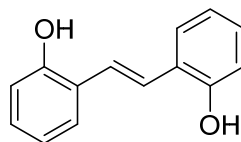
1a

(*E*)-2-(2-Hydroxystyryl)phenol (1a)
White solid, mp 183-185 ºC; lit.[1] mp 188-190 ºC. Spectral data consistent with those reported.[1]

[1] S. K. Nayak, S. Bhatt, K. Roy, , Intramolecular ketone-olefin radical cyclization with low-valent titanium reagent: synthesis of benzopyrans, Synth. Commun. 40 (2010) 2736-2746.
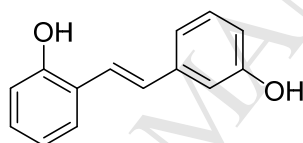
2a

(*E*)-2-(3-Hydroxystyryl)phenol (2a)
White solid, mp 118-120 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.52 (1H, dd, J = 7.8, 1.5 Hz), 7.42 (1H, d, J = 16.5 Hz), 7.14 (1H, t, J = 7.8 Hz), 7.10-7.05 (2H, m), 7.00-6.97 (2H, m), 6.84-6.80 (2H, m), 6.68 (dt, J = 7.8, 1.5 Hz) (distinct OH signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.6, 156.1, 141.0, 125.7 (C) 130.5, 129.5, 129.3, 127.5, 124.8, 120.8, 119.2, 116.6, 115.3, 113.7 (CH); IR ν_{max} (cm^{-1}) 3360 (br, OH); HR ESMS *m/z* 211.0761 (M−H)$^{\ominus}$. Calcd. for C$_{14}$H$_{11}$O$_2$, 211.0759.
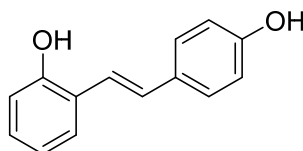
3a

(*E*)-2-(4-Hydroxystyryl)phenol (3a)
White solid, mp 190-192 ºC, lit.[1] mp 192-194 ºC; ^{1}H NMR (500 MHz, DMSO-d$_6$) δ 9.50 (2H, br s, OH), 7.52 (1H, dd, J = 6.5 Hz), 7.37 (2H, d, J = 8.5 Hz), 7.19 (1H, d, J = 16.5 Hz), 7.08 (1H, d, J = 16.5 Hz), 7.04 (1H, td, J = 7.8, 1.5 Hz), 6.84 (1H, dd, J = 7.8, 1 Hz), 6.79 (1H, br t, J ∼ 7.5 Hz),

6.76 (2H, d, J = 8.5 Hz); ^{13}C NMR (125 MHz, DMSO-d_6) δ 157.0, 154.6, 128.8, 124.3 (C) 127.8, 127.7, 127.5 (x 2), 126.0, 120.4, 119.2, 115.7, 115.5 (x 2) (CH); IR ν$_{max}$ (cm^{-1}) 3400 (br, OH); HR ESMS m/z 211.0754 (M−H)$^\ominus$. Calcd. for $C_{14}H_{11}O_2$, 211.0759.

[1]M. A. Ali, K. Kondo, Y. Tsuda, Studies on crude drugs effective on visceral larva migrans .15. synthesis and nematocidal activity of hydroxystilbenes, Chem. Pharm. Bull. 40 (1992) 1130-1136.
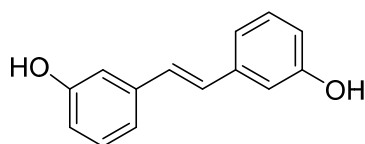
4a

(*E*)-3-(3-Hydroxystyryl)phenol (4a)

White solid, mp 149-151 ºC, lit.[1] mp 152-153 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.15 (2H, t, J = 7.8 Hz), 7.01 (2H, s), 7.00-6.95 (4H, br m), 6.69 (2H, dd, J = 7.8, 1.5 Hz) (distinct OH signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.7 (x 2), 140.2 (x 2) (C) 130.6 (x 2), 129.6 (x 2), 119.3 (x 2), 115.7 (x 2), 113.8 (x 2) (CH); IR ν$_{max}$ (cm^{-1}) 3420 (br, OH); HR ESMS m/z 211.0756 (M−H)$^\ominus$. Calcd. for $C_{14}H_{11}O_2$, 211.0759.

[1]M. A. Ali, K. Kondo, Y. Tsuda, Studies on crude drugs effective on visceral larva migrans .15. synthesis and nematocidal activity of hydroxystilbenes, Chem. Pharm. Bull. 40 (1992) 1130-1136.
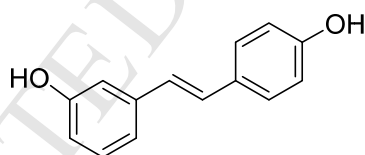
5a

(*E*)-3-(4-Hydroxystyryl)phenol (5a)

White solid, mp 211-213 ºC; lit.[1] mp 216-218 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.35 (2H, br d, J ~ 8 Hz), 7.12 (1H, br t, J ~ 8 Hz), 7.00 (1H, d, J = 16.5 Hz), 6.96 (1H, br d, J ~ 8 Hz), 6.95 (1H, br s), 6.87 (1H, d, J = 16.5 Hz), 6.78 (2H, br d, J ~ 8 Hz), 6.66 (1H, br d, J ~ 8 Hz) (distinct OH signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.6, 158.3, 140.7, 130.4 (C) 130.5, 129.5, 128.8 (x 2), 126.8, 119.0, 116.5 (x 2), 115.1, 113.5 (CH); IR ν$_{max}$ (cm^{-1}) 3350 (br, OH); HR ESMS m/z 211.0757 (M−H)$^\ominus$. Calcd. for $C_{14}H_{11}O_2$, 211.0759.

[1]M. A. Ali, K. Kondo, Y. Tsuda, Studies on crude drugs effective on visceral larva migrans .15. synthesis and nematocidal activity of hydroxystilbenes , Chem. Pharm. Bull. 40 (1992) 1130-1136.
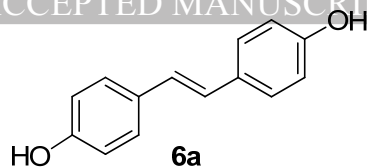
(*E*)-4-(4-Hydroxystyryl)phenol (6a)

Yellowish solid, mp 263-265 ºC; lit.[1] mp 279-281 ºC. Spectral data consistent with those reported:

[1] A. K. Sinha, V. Kumar, A. Sharma, A. Sharma, R. Kumar, An unusual, mild and convenient one-pot two-step access to (E)-stilbenes from hydroxy-substituted benzaldehydes and phenylacetic acids under microwave activation: a new facet of the classical Perkin reaction , Tetrahedon 63 (2007) 11070-11077.
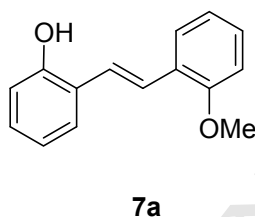
(*E*)-2-(2-Methoxystyryl)phenol (7a)

White solid, mp 104-106 ºC, lit.[1] mp 105-106 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.59 (1H, dd, J = 7.8, 1.5 Hz), 7.53 (1H, dd, J = 7.8, 1.5 Hz), 7.48 (2H, s), 7.15 (1H, td, J = 7.6, 1.5 Hz), 7.06 (1H, td, J = 7.6, 1.5 Hz), 6.90-6.80 (4H, br m), 5.00 (1H, br s, OH), 3.79 (3H, s); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.1, 155.8, 128.2, 126.4 (C) 129.3, 129.2, 127.3, 127.0, 124.8, 123.9, 121.7, 120.8, 116.6, 112.0 (CH) 56.0 (CH$_3$); IR ν_{max} (cm^{-1}) 3500 (br, OH); HR ESMS m/z 225.0913 (M−H)$^{\ominus}$. Calcd. for C$_{15}$H$_{13}$O$_2$, 225.0916.

[1] C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
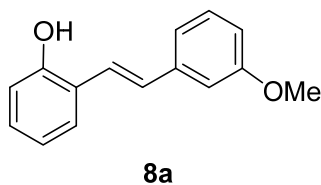
(*E*)-2-(3-Methoxystyryl)phenol (8a)

Yellowish solid, mp 85-87 ºC; lit.[1] mp 89-90 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.51 (1H, br d, J ~ 8 Hz), 7.47 (1H, d, J = 16.2 Hz), 7.20 (1H, t, J = 8 Hz), 7.11 (1H, d, J = 16.2 Hz), 7.10-7.05 (3H, br m), 6.85-6.80 (2H, m), 6.76 (1H, dd, J = 8, 1.5 Hz), 4.90 (1H, br s, OH), 3.77 (3H, s); ^{13}C NMR (125 MHz, CD$_3$OD) δ 161.3, 156.1, 141.0, 125.7 (C), 130.5, 129.6, 129.2, 127.6, 125.1, 120.8, 120.1, 116.6, 114.0, 112.5 (CH), 55.6 (CH$_3$); IR ν_{max} (cm^{-1}) 3400 (br, OH); HR ESMS m/z 227.1073 (M+H)$^{+}$. Calcd. for C$_{15}$H$_{15}$O$_2$, 227.1072.

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
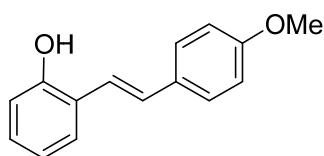
9a

(*E*)-2-(4-Methoxystyryl)phenol (9a)

White solid, mp 134-135 ºC, lit.[1] mp 117-118 ºC, lit.[2] mp 153 ºC. Spectral data consistent with those reported.[1,2]

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
[2]M. Chalal, D. Vervandier-Fasseur, P. Meunier, H. Cattey, J.-C. Hierso, Syntheses of polyfunctionalized resveratrol derivatives using Wittig and Heck protocols, Tetrahedron 68 (2012) 3899-3907.
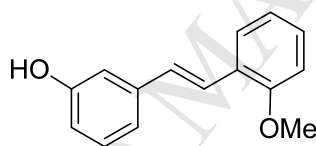
10a

(*E*)-3-(2-Methoxystyryl)phenol (10a)

White solid, mp 72-73 ºC, lit.[1] mp 78-79 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.52 (1H, dd, J = 7.4, 1.5 Hz), 7.42 (1H, d, J = 16.1 Hz), 7.16 (1H, td, J = 8, 1.5 Hz), 7.13 (1H, t, J = 7.8 Hz), 7.02 (1H, d, = 16.1 Hz), 7.00-6.95 (2H, m), 6.90-6.85 (2H, m), 6.70 (1H, dd, J = 8.2 Hz, 2.5 Hz), 4.80 (1H, br s, OH), 3.77 (3H, s); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.5, 158.2, 140.8, 124.4 (C) 130.6, 129.8, 129.7, 127.3, 127.2, 121.7, 119.3, 115.5, 113.7, 112.1 (CH), 56.0 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3400 (br, OH); HR ESMS *m/z* 225.0920 (M−H)$^{\ominus}$. Calcd. for C$_{15}$H$_{13}$O$_2$, 225.0916.

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
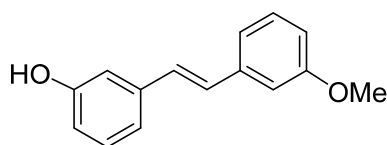
11a

(*E*)-3-(3-Methoxystyryl)phenol (11a)

White solid, mp 78-80 ºC, lit.[1] mp 74-75 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.19 (1H, t, J = 7.8 Hz), 7.14 (1H, t, J = 7.8 Hz), 7.05-6.95 (5H, br m), 6.76 (1H, dd, J = 8.2, 2.5 Hz), 6.72 (1H, dd, J = 7.8, 2 Hz), 3.75 (3H, s) (a distinct OH signal not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 161.3, 158.6, 140.1 (x 2) (C), 130.6, 130.5, 129.9, 129.5, 120.1, 119.4, 115.7, 114.2, 113.9, 112.7 (CH), 55.6 (CH$_3$); IR ν$_{max}$ 3400 (br, OH) cm^{-1}; HR ESMS m/z 225.0919 (M−H)$^\ominus$. Calcd. for C$_{15}$H$_{13}$O$_2$, 225.0916.

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
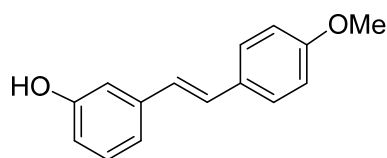
12a

(*E*)-3-(4-Methoxystyryl)phenol (12a)
White solid, mp 162-164 ºC, lit.[1] mp 159-160 ºC, lit.[2] mp 151 ºC. Spectral data consistent with those reported.[1,2]

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
[2]M. Chalal, D. Vervandier-Fasseur, P. Meunier, H. Cattey, J.-C. Hierso, Syntheses of polyfunctionalized resveratrol derivatives using Wittig and Heck protocols, Tetrahedron 68 (2012) 3899-3907.
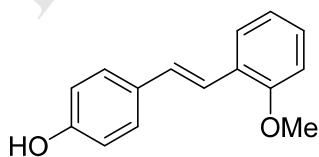
13a

(*E*)-4-(2-Methoxystyryl)phenol (13a)
Yellowish solid, mp 119-121 ºC, lit.[1] mp 145-146 ºC. Spectral data consistent with those reported.[1,2]

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
[2]T. Song, B. Zhou, G.-W. Peng, Q.-B. Zhang, L.-Z. Wu, Q. Liu, W. Yong, Aerobic Oxidative Coupling of Resveratrol and its Analogues by Visible Light Using Mesoporous Graphitic Carbon Nitride (mpg-C3N4) as a Bioinspired Catalyst, Chem. Eur. J. 20 (2014) 678-682.

14a
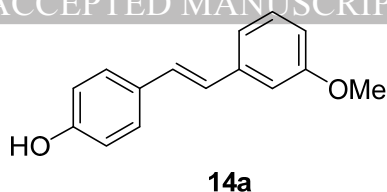
(*E*)-4-(3-Methoxystyryl)phenol (14a)

White solid, mp 125-127 ºC, lit.[1] mp 118-119 ºC, lit.[2] mp 132-134 ºC. Spectral data consistent with those reported.[1,2]

[1]C. J. Lion, C. S. Matthews, M. F. G. Stevens, A. D. Westwell, Synthesis, antitumor evaluation, and apoptosis-inducing activity of hydroxylated (E)-stilbenes , J. Med. Chem. 48 (2005) 1292-1295.
[2]A. K. Sinha, V. Kumar, A. Sharma, A. Sharma, R. Kumar, An unusual, mild and convenient one-pot two-step access to (E)-stilbenes from hydroxy-substituted benzaldehydes and phenylacetic acids under microwave activation: a new facet of the classical Perkin reaction , Tetrahedon 63 (2007) 11070-11077.

15a
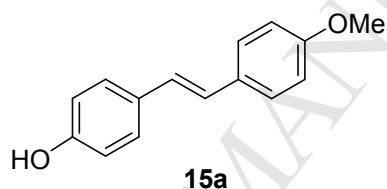
(*E*)-4-(4-Methoxystyryl)phenol (15a)

Yellowish solid, mp 198-200 ºC, lit.[1] mp 211-212 ºC. Spectral data consistent with those reported.[1]

[1]M. Chalal, D. Vervandier-Fasseur, P. Meunier, H. Cattey, J.-C. Hierso, Syntheses of polyfunctionalized resveratrol derivatives using Wittig and Heck protocols, Tetrahedron 68 (2012) 3899-3907.

1b
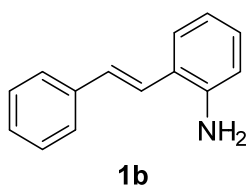
(*E*)-2-Styrylaniline (1b)

Yellowish solid, mp 102-104 ºC. Spectral data consistent with those reported: M. Shen, B. E. Leslie, T. G. Driver, Dirhodium(II)-catalyzed intramolecular C-H amination of aryl azides, Angew. Chem. Int. Ed. 47 (2008) 5056-5059.

2b
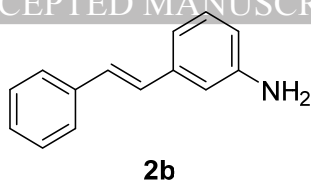
(*E*)-3-Styrylaniline (2b)

Yellowish solid, mp 109-111 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.57 (2H, d, *J* = 7.8 Hz), 7.42 (2H, t, *J* = 7.5 Hz), 7.33 (1H, td, *J* = 7.6, 1 Hz), 7.20 (1H, t, *J* = 7.8 Hz), 7.10 (2H, AB system, *J* = 16.2 Hz), 7.00 (1H, d, *J* = 7.6 Hz), 6.88 (1H, br s), 6.65 (1H, dt, *J* = 7.8, 1 Hz), 3.70 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 146.6, 138.3, 137.3 (C), 129.5, 128.8, 128.6 (x 2), 128.4, 127.5, 126.4 (x 2), 117.2, 114.6, 112.8 (CH).

^{1}H NMR data consistent with those reported in: K. Motoshima, T. Noguchi-Yachide, K. Sugita, Y. Hashimoto, M. Ishikawa, Separation of alpha-glucosidase-inhibitory and liver X receptor-antagonistic activities of phenethylphenyl phthalimide analogs and generation of LXR alpha-selective antagonists, Bioorg. Med. Chem. 17 (2009) 5001-5014.

3b
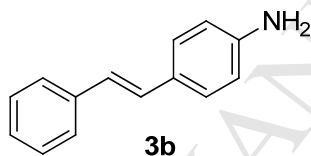
(*E*)-4-styrylaniline (3b)

Yellowish solid, mp 147-149 ºC. Spectral data consistent with those reported in: R. K. Arvela, N. E. Leadbeater, M. S. Sangi, V. A. Williams, P. Granados, R. D. Singer, A reassessment of the transition-metal free Suzuki-type coupling methodology, J. Org. Chem. 70 (2005) 161-168.

4b
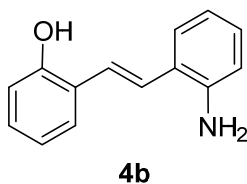
(*E*)-2-(2-Aminostyryl)phenol (4b)

White solid, mp 155-157 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.59 (1H, dd, *J* = 7.8, 1.5 Hz), 7.43 (1H, dd, *J* = 7.8, 1 Hz), 7.30 (2H, AB system, *J* = 16 Hz), 7.06 (1H, dt, *J* = 8.2, 1.7 Hz), 7.01 (1H, dt, *J* = 7.8, 1.5 Hz), 6.85-6.80 (2H, m), 6.75-6.70 (2H, m), 4.80 (3H, br s, OH, NH$_2$); ^{13}C NMR (125 MHz, CD$_3$OD) δ 156.0, 145.9, 126.4, 125.8 (C) 129.2, 129.1, 127.6, 127.3, 125.3, 124.8, 120.8, 119.8, 117.7, 116.6 (CH); IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH), 3400, 3320 (NH$_2$); HR ESMS *m/z* 212.1078 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

5b
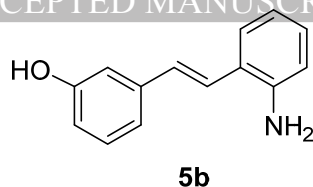
(*E*)-3-(2-Aminostyryl)phenol (5b)

Yellowish solid, mp 68-70 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.40 (1H, dd, *J* = 7.8, 1 Hz), 7.26 (1H, d, *J* = 16.2 Hz), 7.14 (1H, t, *J* = 7.8 Hz), 7.05-7.00 (3H, m), 6.91 (1H, d, *J* = 16.2 Hz), 6.75-6.68 (3H, br m) (distinct OH/NH$_2$ signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.6, 146.1, 140.7, 127.3 (C) 130.6, 130.1, 129.5, 125.2, 124.9, 119.8, 119.3, 117.8, 115.4, 113.9 (CH); IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH, NH$_2$); HR ESMS *m/z* 212.1073 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

6b
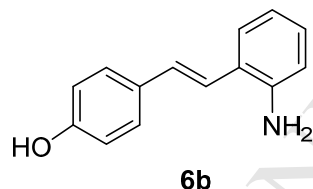
(*E*)-4-(2-Aminostyryl)phenol (6b)

Yellowish solid, mp 193-195 ºC; ^{1}H NMR (500 MHz, DMSO-d$_6$) δ 9.40 (1H, br s, OH), 7.44 (2H, d, *J* = 8.5 Hz), 7.36 (1H, dd, *J* = 7.8, 1.5 Hz), 7.15 (1H, d, *J* = 16.1 Hz), 6.92 (1H, td, *J* = 7.8, 1.5 Hz), 6.87 (1H, d, *J* = 16.1 Hz), 6.75 (2H, d, *J* = 8.5 Hz), 6.63 (1H, dd, *J* = 7.8, 1 Hz), 6.53 (1H, br t, *J* ~ 8 Hz), 5.20 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, DMSO-d$_6$) δ 156.8, 145.9, 129.0, 121.5 (C) 127.8, 127.7 (x 2), 127.0, 125.1, 121.2, 116.3, 115.4, 115.3 (x 2) (CH); IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH, NH$_2$); HR ESMS *m/z* 212.1073 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

7b
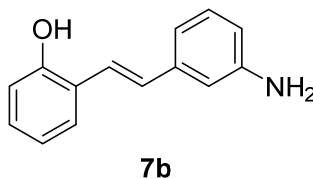
(*E*)-2-(3-Aminostyryl)phenol (7b)

White solid, mp 159-161 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.51 (1H, d, *J* = 7.5 Hz), 7.41 (1H, d, *J* = 16.1 Hz), 7.10-7.00 (3H, m), 6.93 (1H, br s), 6.88 (1H, br d, *J* ~ 7.8 Hz), 6.80 (2H, m), 6.62 (1H, dt, *J* = 7.8, 1 Hz), (distinct OH, NH$_2$ signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ (C) 156.1, 148.7, 140.4, 125.9 (CH) 130.2, 129.7, 129.3, 127.4, 124.3, 120.8, 118.1, 116.6, 115.9, 114.3; IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH), 3350, 3290 (NH$_2$); HR ESMS *m/z* 212.1074 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

8b
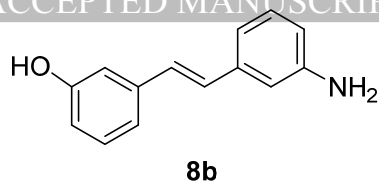
(*E*)-3-(3-Aminostyryl)phenol (8b)

White solid, mp 165-167 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.15 (1H, t, *J* = 7.8 Hz), 7.08 (1H, t, *J* = 7.8 Hz), 7.00-6.90 (5H, m), 6.87 (1H, br d, *J* = 7.8 Hz), 6.68 (1H, dd, *J* = 7.8, 1.6 Hz), 6.63 (1H, dd, *J* = 7.8, 1.6 Hz), (distinct OH, NH$_2$ signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 158.6, 148.7, 140.2, 139.5 (C) 130.5, 130.3, 130.0, 129.2, 119.2, 118.0, 116.2, 115.6, 114.3, 113.8 (CH); IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH), 3300, 3180 (NH$_2$); HR ESMS *m/z* 212.1075 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

9b
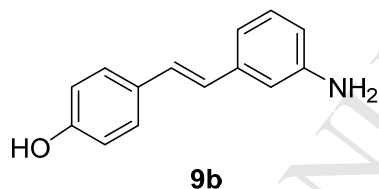
(*E*)-4-(3-Aminostyryl)phenol (9b)

Yellowish solid, mp 241-243 ºC; ^{1}H NMR (500 MHz, DMSO-d$_6$) δ 7.40 (2H, d, *J* = 8.5 Hz), 7.01 (1H, t, *J* = 7.8 Hz), 6.96 (1H, d, *J* = 16.1 Hz), 6.86 (1H, d, *J* = 16.1 Hz), 6.78 (2H, d, *J* = 8.5 Hz), 6.74 (1H, t, *J* = 1.5 Hz), 6.71 (1H, br d, *J* ~ 7.5 Hz), 6.47 (1H, dt, *J* = 7.8, 1.5 Hz), 5.00 (3H, br s, OH, NH$_2$); ^{13}C NMR (125 MHz, DMSO-d$_6$) δ 157.1, 148.7, 137.9, 128.2 (C) 129.0, 127.7 (x 2), 127.3, 126.0, 115.5 (x 2), 114.3, 113.2, 111.3 (CH); IR ν$_{max}$ (cm^{-1}) 3500-3200 (br, OH, NH$_2$); HR ESMS *m/z* 212.1076 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.

10b
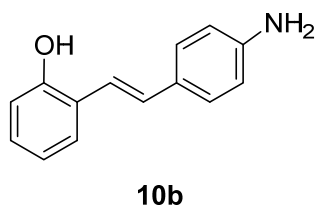
(*E*)-2-(4-Aminostyryl)phenol (10b)

Yellowish solid, mp 179-181 ºC; ^{1}H NMR (500 MHz, CD$_3$OD) δ 7.48 (1H, d, *J* = 7.8 Hz), 7.30 (2H, d, *J* = 7.8 Hz), 7.24 (1H, d, *J* = 16.1 Hz), 7.05-7.00 (2H, m), 6.82-6.79 (2H, m), 6.70 (2H, d, *J* = 7.8 Hz), (distinct OH, NH$_2$ signals not detected, perhaps merged with the solvent signal); ^{13}C NMR (125 MHz, CD$_3$OD) δ 155.6, 148.3, 129.8, 126.5 (C) 129.6, 128.6, 128.4 (x 2), 127.0, 120.8, 120.7, 116.6 (x 3); IR ν$_{max}$ (cm^{-1}) 3500-3100 (br, OH, NH$_2$); HR ESMS *m/z* 212.1073 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.
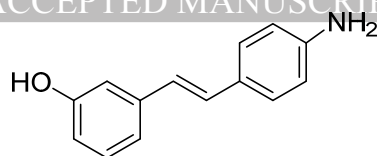
11b

(*E*)-3-(4-Aminostyryl)phenol (11b)

Yellowish solid, mp 146-148 ºC; ^{1}H NMR (500 MHz, DMSO-d$_6$) δ 9.30 (1H, br s, OH), 7.26 (2H, d, *J* = 8.5 Hz), 7.11 (1H, t, *J* = 7.8 Hz), 6.95 (1H, d, *J* = 16.1 Hz), 6.92 (1H, br d, *J* ~ 8 Hz), 6.88 (1H, t, *J* = 2 Hz), 6.80 (1H, d, *J* = 16.1 Hz), 6.60 (1H, dt, *J* = 7.8, 1.5 Hz), 6.56 (2H, d, *J* = 8.5 Hz), 5.30 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, DMSO-d$_6$) δ 157.5, 148.7, 139.3, 124.7 (C) 129.4, 128.8, 127.6 (x 2), 123.0, 116.8, 113.9 (x 2), 113.7, 112.3 (CH); IR ν$_{max}$ (cm^{-1}) 3500-32 00 (br, OH, NH$_2$); HR ESMS *m/z* 212.1079 (M+H)$^+$. Calcd. for C$_{14}$H$_{14}$NO, 212.1075.
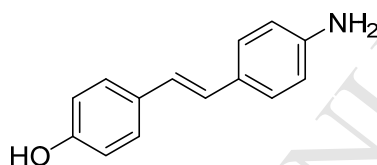
12b

(*E*)-4-(4-Aminostyryl)phenol (12b)

Yellowish solid, mp 82-84 ºC. Spectral data consistent with those reported in: J. Y. Kim, J. W. Lee, Y. S. Kim, Y. Lee, Y. B. Ryu, S. Kim, H. W. Ryu, M. J. Curtis-Long, K. W. Lee, W. S. Lee, K. H. Park, A Novel competitive class of alpha-glucosidase inhibitors: (E)-1-phenyl-3-(4-styrylphenyl)urea derivatives, ChemBioChem 11 (2010) 2125-2131.
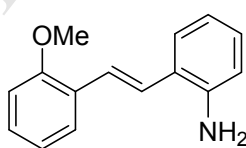
13b

(*E*)-2-(2-Methoxystyryl)aniline (13b)

Yellowish solid, mp 101-103 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.56 (1H, dd, *J* = 7.8, 1.5 Hz), 7.42 (1H, dd, *J* = 7.8, 1 Hz), 7.33 (1H, d, *J* = 16.1 Hz), 7.23 (1H, td, *J* = 7.8 Hz, 1.6 Hz), 7.16 (1H, d, *J* = 16.1 Hz), 7.08 (1H, td, *J* = 7.6, 1.5 Hz), 6.96 (1H, br t, *J* = 7.6 Hz), 6.89 (1H, br d, *J* = 7.8 Hz), 6.80 (1H, br t, *J* = 7.6 Hz), 6.70 (1H, dd, *J* = 7.8, 1 Hz), 3.90 (2H, br s, NH$_2$), 3.87 (3H, s); ^{13}C NMR (125 MHz, CDCl$_3$) δ 156.9, 143.8, 126.8, 124.6 (C), 128.6, 128.4, 127.3, 126.6, 125.3, 124.7, 120.7, 119.1, 116.2, 110.9 (CH), 55.5 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3385, 3360 (NH$_2$); HR ESMS *m/z* 226.1231 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
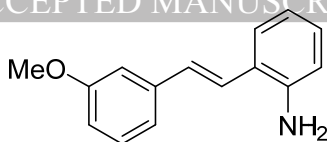
14b

(*E*)-2-(3-Methoxystyryl)aniline (14b)

Yellowish oil; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.46 (1H, br d, *J* ~ 8 Hz), 7.34 (1H, t, *J* = 8 Hz), 7.22 (1H, d, *J* = 16.1 Hz), 7.20-7.15 (2H, m), 7.12 (1H, br s), 7.03 (1H, d, *J* = 16.1 Hz), 6.90-6,85 (2H, br m), 6.75 (1H, br d, *J* ~ 8 Hz), 3.88 (3H, s), 3.75 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 159.8, 144.0, 139.0, 123.5 (C), 129.8, 129.5, 128.6, 127.1, 124.4, 119.0, 118.9, 116.1, 113.0, 111.6 (CH), 55.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3440, 3370 (NH$_2$); HR ESMS *m/z* 226.1229 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
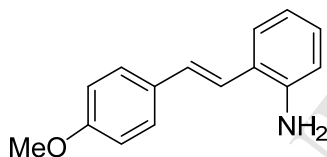
15b

(*E*)-2-(4-Methoxystyryl)aniline (15b)

Yellowish solid, mp 120-122 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.49 (2H, d, *J* = 8 Hz), 7.43 (1H, d, *J* = 7.8 Hz), 7.13 (1H, td, *J* = 7.8, 1 Hz), 7.06 (1H, d, *J* = 16.1 Hz), 6.99 (1H, d, *J* = 16.1 Hz), 6.94 (2H, d, *J* = 8 Hz), 6.85 (1H, br t, *J* = 7.6 Hz), 6.74 (1H, d, *J* = 7.8 Hz), 3.86 (3H, s), 3.70 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 159.2, 143.2, 130.3, 124.4 (C), 129.9, 128.2, 127.6 (x 2), 126.9, 122.0, 119.4, 116.4, 114.1 (x 2) (CH), 55.2 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3440, 3360 (NH$_2$); HR ESMS *m/z* 226.1236 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
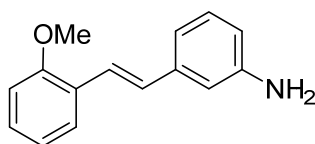
16b

(*E*)-3-(2-Methoxystyryl)aniline (16b)

Yellowish oil; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.59 (1H, dd, *J* = 7.5, 1.5 Hz), 7.44 (1H, d, *J* = 16.5 Hz), 7.25 (1H, td, *J* = 7.8, 1.5 Hz), 7.15 (1H, t, *J* = 7.8 Hz), 7.03 (1H, d, *J* = 16.5 Hz), 7.00-6.90 (4H, br m), 6.60 (1H, dd, *J* = 7.8, 1.5 Hz), 3.88 (3H, s), 3.60 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 156.9, 146.6, 139.1, 126.5 (C), 129.5, 129.2, 128.6, 126.4, 123.3, 120.7, 117.5, 114.5, 113.0, 111.0 (CH), 55.6 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3450, 3370 (NH$_2$); HR ESMS *m/z* 226.1227 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
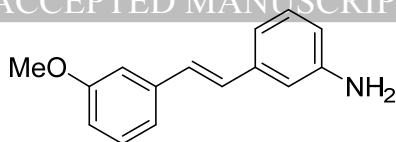
17b

(*E*)-3-(3-Methoxystyryl)aniline (17b)

White solid, mp 75-77 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.31 (1H, t, *J* = 7.8 Hz), 7.20 (1H, t, *J* = 7.8 Hz), 7.15 (1H, d, *J* = 7.8 Hz), 7.08 (3H, m), 6.97 (1H, d, *J* = 7.8 Hz), 6.87 (2H, m), 6.64 (1H, br d, *J* ~ 8 Hz), 3.88 (3H, s), 3.70 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 159.8, 146.6, 138.8, 138.1 (C) 129.5, 129.4, 129.1, 128.3, 119.1, 117.2, 114.6, 113.1, 112.8, 111.7 (CH) 55.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3450, 3370 (NH$_2$); HR ESMS *m/z* 226.1233 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
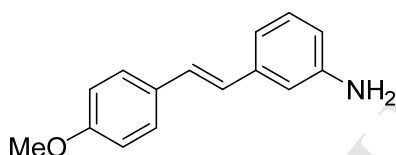
18b

(*E*)-3-(4-Methoxystyryl)aniline (18b)

White solid, mp 143-145 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.46 (2H, d, *J* = 8.5 Hz), 7.16 (1H, d, *J* = 7.8 Hz), 7.04 (1H, d, *J* = 16.2 Hz), 6.95-6.90 (4H, m), 6.84 (1H, t, *J* = 1 Hz), 6.60 (1H, dd, *J* = 7.8, 1.5 Hz), 3.85 (3H, s), 3.70 (2H, br s, NH$_2$); ^{13}C NMR (125 MHz, CDCl$_3$) δ 159.2, 146.6, 138.7, 130.2 (C), 129.5, 128.0, 127.6 (x 2), 126.8, 117.0, 114.3, 114.1 (x 2), 112.7 (CH), 55.2 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3400, 3330 (NH$_2$); HR ESMS *m/z* 226.1230 (M+H)$^+$. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.
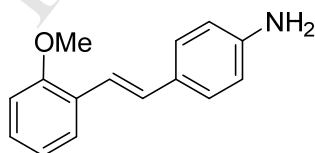
19b

(*E*)-4-(2-Methoxystyryl)aniline (19b)

Yellowish solid, mp 70-72 ºC, lit.[1] mp 68-70 ºC; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.57 (1H, dd, *J* = 7.8, 1 Hz), 7.37 (2H, d, *J* = 8.5 Hz), 7.30 (1H, d, *J* = 16.2 Hz), 7.21 (1H, br t, *J* ~ 78 Hz), 7.03 (1H, d, *J* = 16.2 Hz), 6.96 (1H, t, *J* = 7.5 Hz), 6.89 (1H, d, *J* = 7.8 Hz), 6.71 (2H, d, *J* = 8.5 Hz), 3.88 (3H, s) (a distinct NH$_2$ signal not detected); ^{13}C NMR (125 MHz, CDCl$_3$) δ 156.7, 145.9, 130.0, 128.8 (C), 129.1, 127.9, 127.8 (x 2), 126.0, 120.7, 119.9, 115.2 (x 2), 110.9 (CH), 55.5 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3460, 3370 (NH$_2$); HR ESMS *m/z* (M+H)$^+$ 226.1230. Calcd. for C$_{15}$H$_{16}$NO, 226.1232.

[1]H. Ulrich, D. V. Rao, F. A. Stuber, A. A. R. Sayigh, Photodimerization of substituted stilbenes, J. Org. Chem. 35 (1970) 1121-1125.
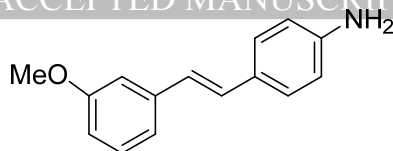
20b

(*E*)-4-(3-Methoxystyryl)aniline (20b)

Yellowish solid, mp 76-78 ºC, lit.[1] mp 70-71 ºC. Spectral data consistent with those reported.[1]

[1]B. Sun, J. Hoshino, K. Jermihov, L. Marler, J. M. Pezzuto, A. D. Mesecar, M. Cushman, Design, synthesis, and biological evaluation of resveratrol analogues as aromatase and quinone reductase 2 inhibitors for chemoprevention of cancer, Bioorg. Med. Chem. 18 (2010) 5352-5366.
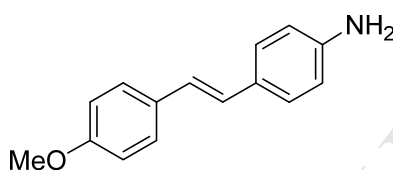
21b

(*E*)-4-(4-Methoxystyryl)aniline (21b)

Yellowish solid, mp 157-159 ºC. Spectral data consistent with those reported in: C. M. Kormos, N. E. Leadbeater, Preparation of nonsymmetrically substituted stilbenes in a one-pot two-step Heck strategy using ethene as a reagent, J. Org. Chem. 73 (2008) 3854-3858.
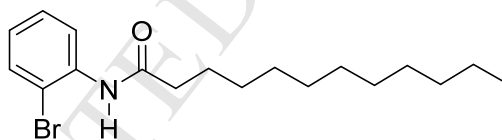
N-(2-Bromophenyl)dodecanamide

Prepared by means of acylation of 2-bromoaniline under the same conditions described in page S-4 for the conversion of **2b** into **23b** (60% yield).

White solid, mp 73-75 °C; ^{1}H NMR (500 MHz, CDCl$_3$) δ 8.35 (1H, br d, *J* ~ 7.8 Hz), 7.60 (1H, br s, NH), 7.51 (1H, dd, *J* = 8, 1.5 Hz), 7.28 (1H, td, *J* = 7.5, 1 Hz), 6.95 (1H, td, *J* = 7.8, 1.5 Hz), 2.41 (2H, t, *J* = 7.5 Hz), 1.74 (2H, quint, *J* = 7.5 Jz), 1.40-1.20 (16H, br m), 0.89 (3H, t, *J* = 7 Hz); ^{13}C NMR (125 MHz, CDCl$_3$) δ 171.3, 122.0, 113.3 (C), 135.8, 132.1, 128.3, 125.0 (CH), 38.0, 31.9, 29.6 (x 2), 29.5, 29.4, 29.3, 29.2, 25.5, 22.7 (CH$_2$), 14.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3280 (NH), 1664 (C=O); HR ESMS *m/z* 354.1432 (M+H)$^+$. Calcd. for C$_{18}$H$_{29}$NO^{79}Br, 354.1433.

S-17
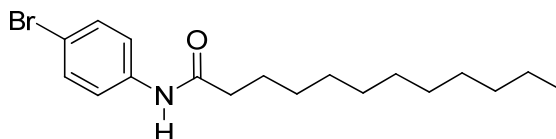
N-(4-bromophenyl)dodecanamide

Prepared by means of acylation of 4-bromoaniline under the same conditions described in page S-4 for the conversion of **2b** into **23b** (62% yield).

Yellowish solid, mp 106-108 °C; lit.[1] mp 106 ºC. Spectral data consistent with those reported.[2]

[1]D. F. Houston, Melting points of the p-bromoanilides of solid fatty acids, J. Am. Chem. Soc. 62 (1940) 1303-1304.

[2]Y. Chen, B. Zhu, F. Zhang, Y. Han, Z. S. Bo, Hierarchical supramolecular self-assembly of nanotubes and layered sheets, Angew. Chem. Int. Ed. 47 (2008) 6015-6018.

22b
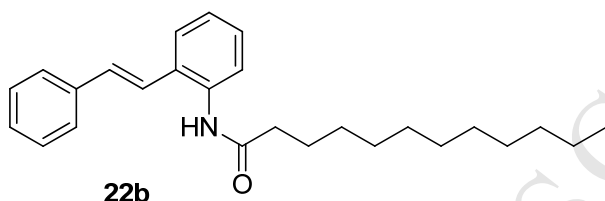
(*E*)-*N*-(2-Styrylphenyl)dodecanamide (22b)

Yellowish solid, mp 100-102 °C; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.80 (1H, br d, *J* ~ 7.5 Hz), 7.51 (1H, br d; *J* ~ 8 Hz), 7.49 (2H, br d, *J* ~ 7.5 Hz), 7.37 (2H, t, *J* = 7.8 Hz), 7.30-7.20 (4H, br m), 6.98 (1H, d, *J* = 16 Hz), 2.40 (2H, t, *J* = 6.8 Hz), 1.75 (2H, m), 1.40-1.20 (16H, br m), 0.88 (3H, t, *J* = 6.8 Hz) (a distinct NH signal not detected); ^{13}C NMR (125 MHz, CDCl$_3$) δ 171.6, 137.0, 132.4, 130.3 (C), 134.8, 128.8 (x 3), 128.3, 128.1, 126.8, 126.6, 125.4, 124.2, 123.5 (CH), 37.6, 31.9, 29.7, 29.6 (x 2), 29.5, 29.4, 29.3, 25.9, 22.7 (CH$_2$), 14.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3285 (NH), 1664 (C=O); HR ESMS *m/z* 377.2796 (M+H)$^+$. Calcd. for C$_{26}$H$_{36}$NO, 378.2797.

23b
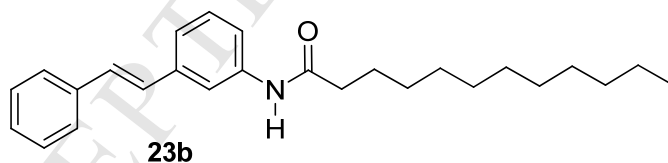
(*E*)-*N*-(3-Styrylphenyl)dodecanamide (23b)

Yellowish solid, mp 103-105 °C; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.80 (1H, br s, NH), 7.48 (2H, br d, *J* = 7.4 Hz), 7.35-7.20 (7H, br m), 7.08 (2H, AB system, *J* = 16.2 Hz), 2.36 (2H, t, *J* = 7.5 Hz), 1.74 (2H, quintuplet, *J* = 7.5 Hz), 1.40-1.20 (16H, br m), 0.88 (3H, t, *J* = 7 Hz); ^{13}C NMR (125 MHz, CDCl$_3$) δ 171.6, 138.4, 138.3, 137.1 (C), 129.3, 128.7 (x 2), 128.3, 127.7, 126.5 (x 2), 122.5, 118.9, 117.7, 117.6 (CH), 37.9, 31.9, 29.6 (x 2), 29.5, 29.4, 29.3, 29.2, 25.6, 22.7 (CH$_2$), 14.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3400 (NH), 1660 (C=O); HR ESMS *m/z* 377.2794 (M+H)$^+$. Calcd. for C$_{26}$H$_{36}$NO, 378.2797.

24b
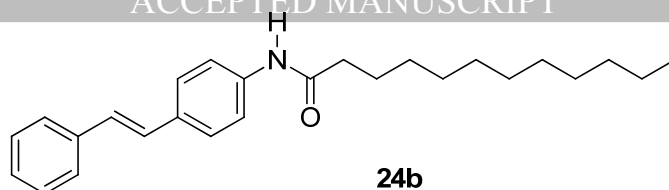
(*E*)-*N*-(4-Styrylphenyl)dodecanamide (24b)

Yellowish solid, mp 168-170 °C; ^{1}H NMR (500 MHz, CDCl$_3$) δ 7.55-7.45 (6H, br m), 7.35 (2H, t, *J* = 7.8 Hz), 7.25 (1H, t, *J* =7.8 Hz), 7.14 (1H, br s, NH), 7.05 (2H, AB system, *J* = 16.2 Hz), 2.36 (2H, t, *J* = 7.5 Hz), 1.73 (2H, quintuplet, *J* = 7.5 Hz), 1.40-1.20 (16H, br m), 0.88 (3H, t, *J* = 6.8 Hz); ^{13}C NMR (125 MHz, CDCl$_3$) δ 171.3, 137.4 (x 2), 128.8 (C) 128.6 (x 2), 128.0, 127.8, 127.4, 127.1 (x 2), 126.4 (x 2), 119.8 (x 2) (CH), 37.9, 31.9, 29.6 (x 2), 29.5, 29.4, 29.3 (x 2), 25.6, 22.7 (CH$_2$), 14.1 (CH$_3$); IR ν$_{max}$ (cm^{-1}) 3300 (NH), 1659 (C=O); HR ESMS *m/z* 377.2798 (M+H)$^{+}$. Calcd. for C$_{26}$H$_{36}$NO, 378.2797.

NMR spectra of stilbene derivatives
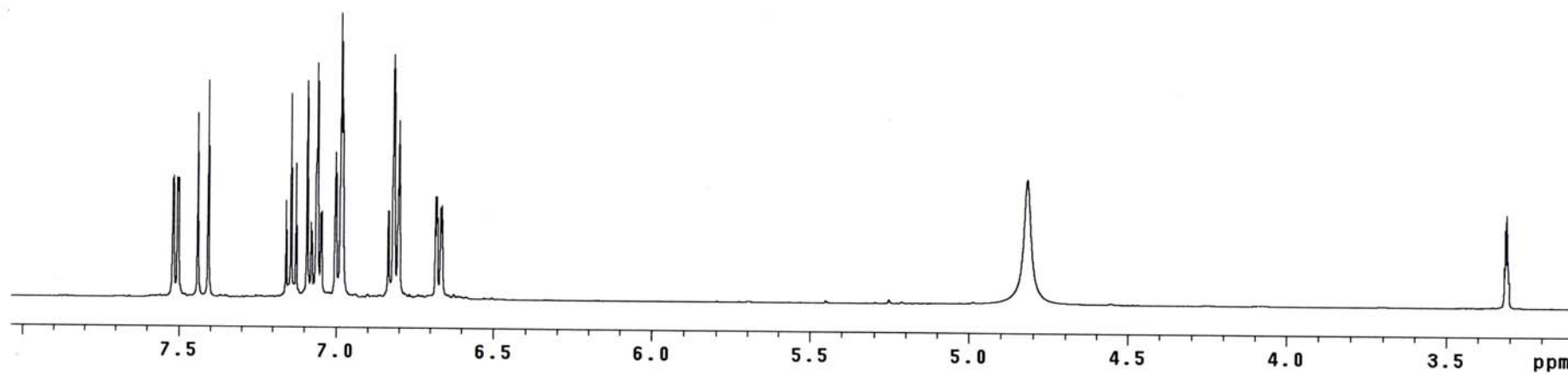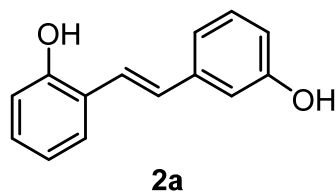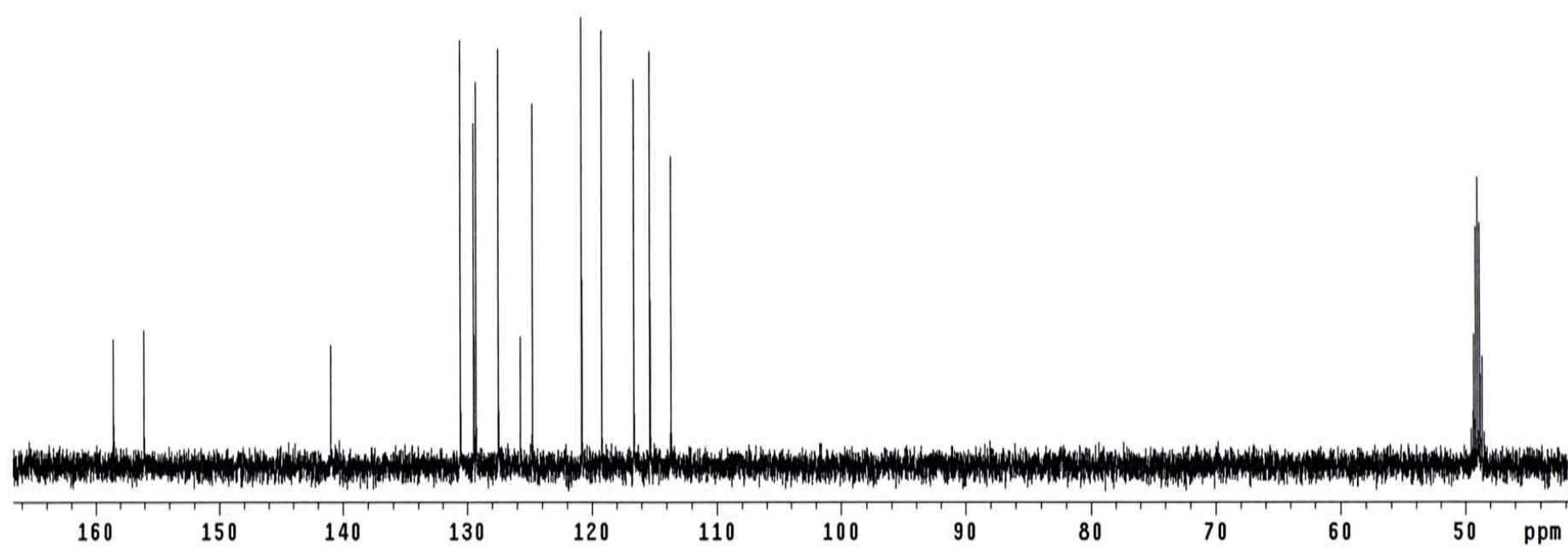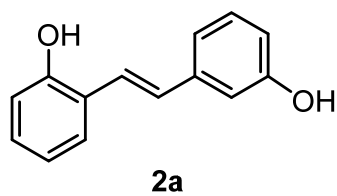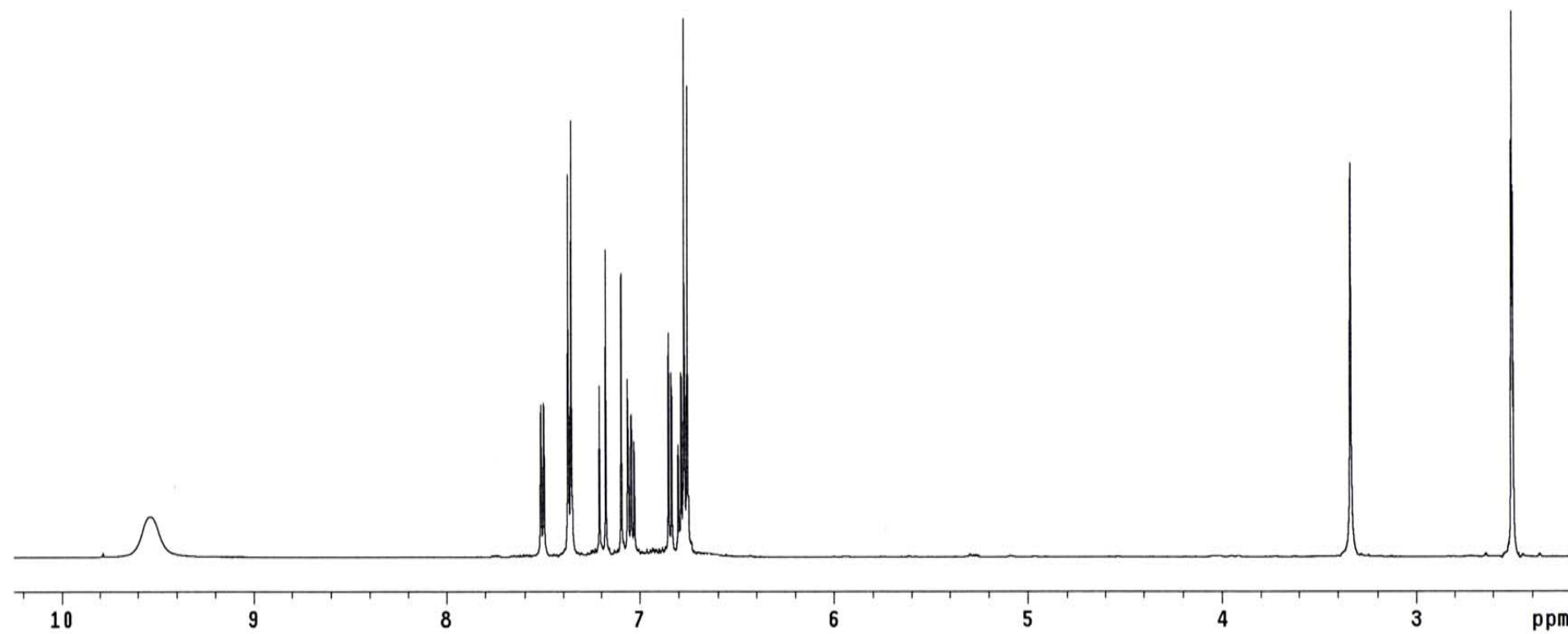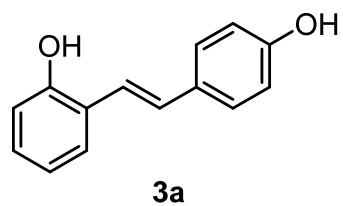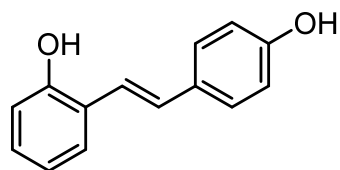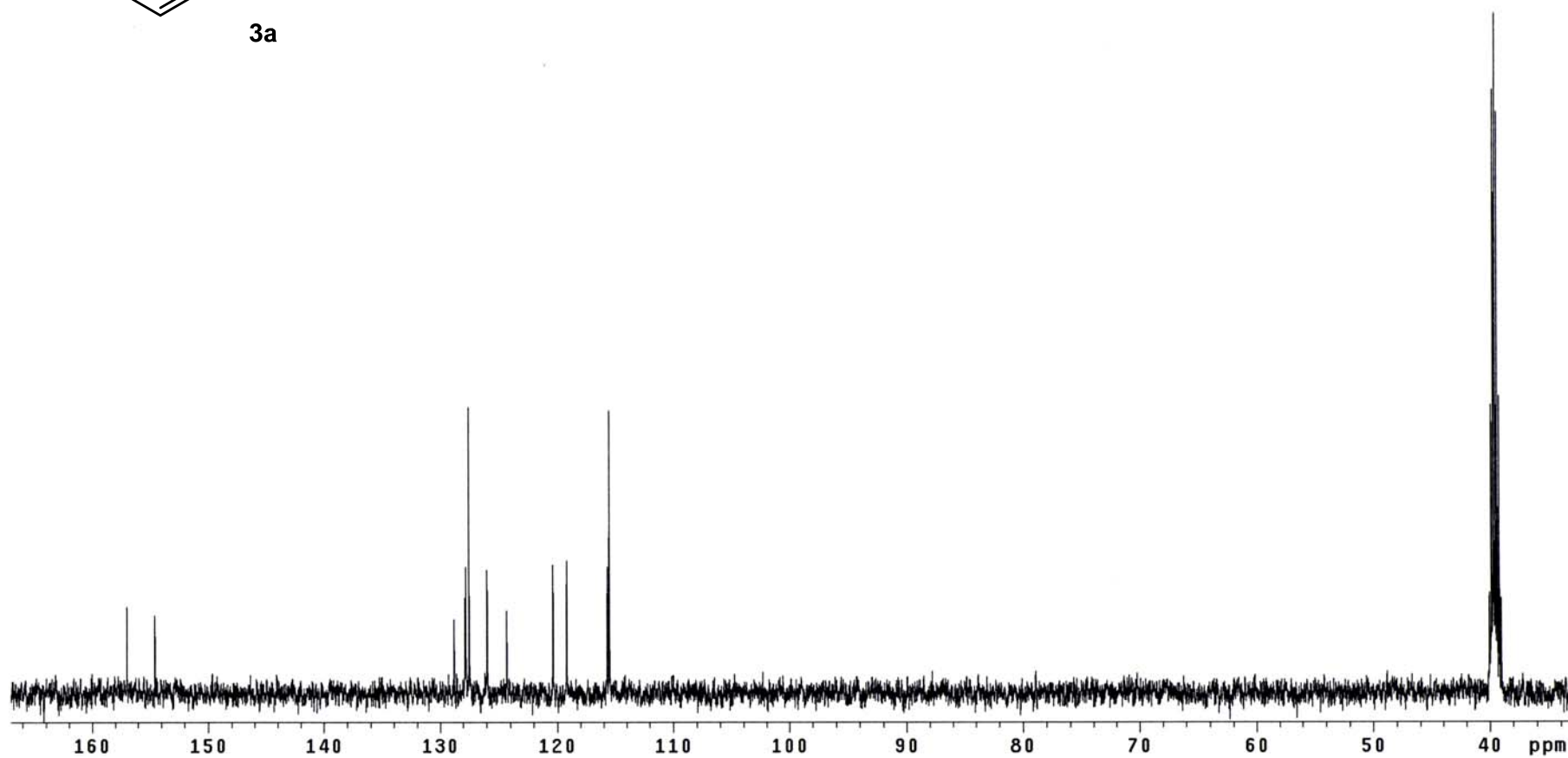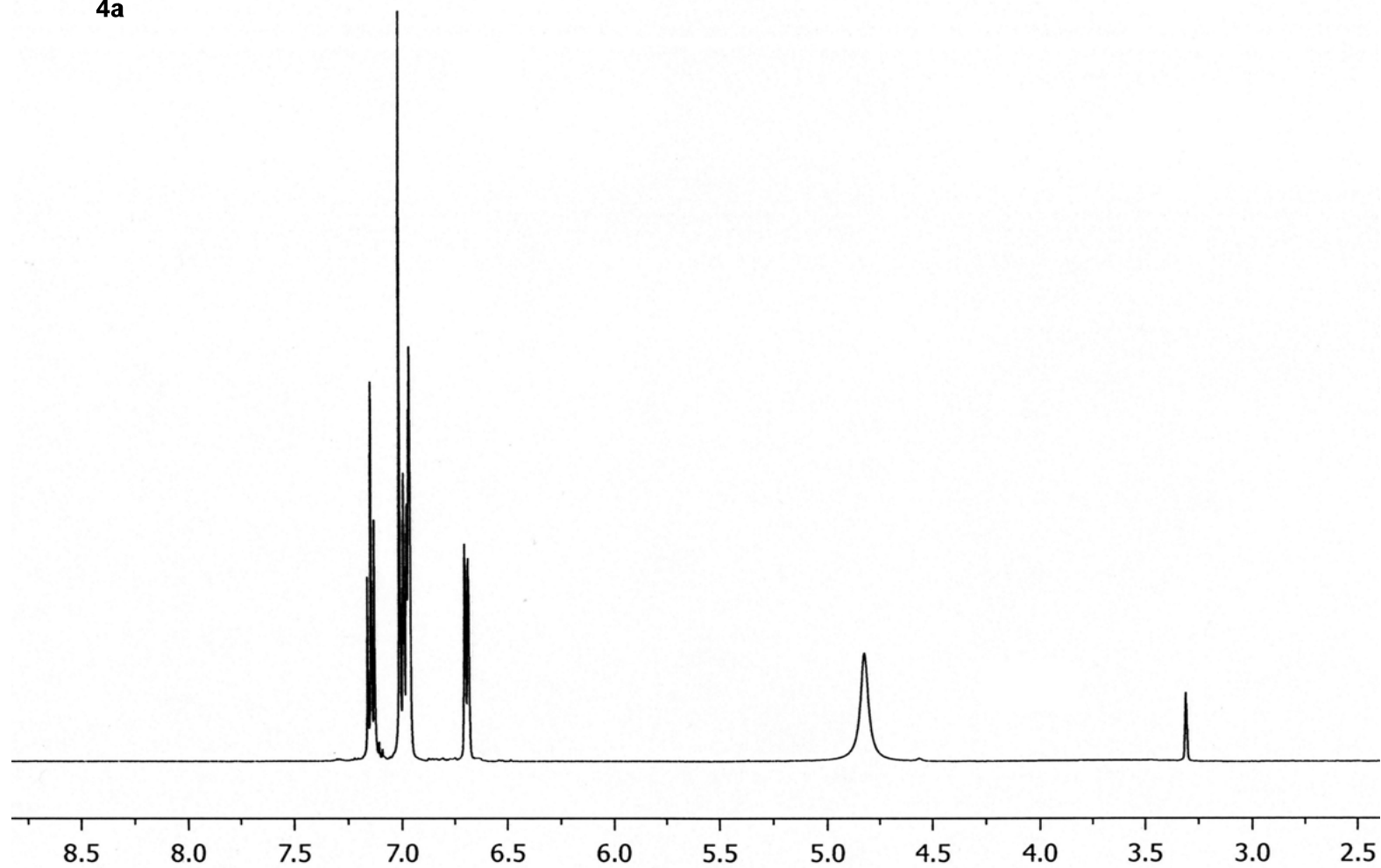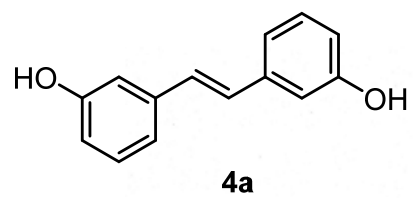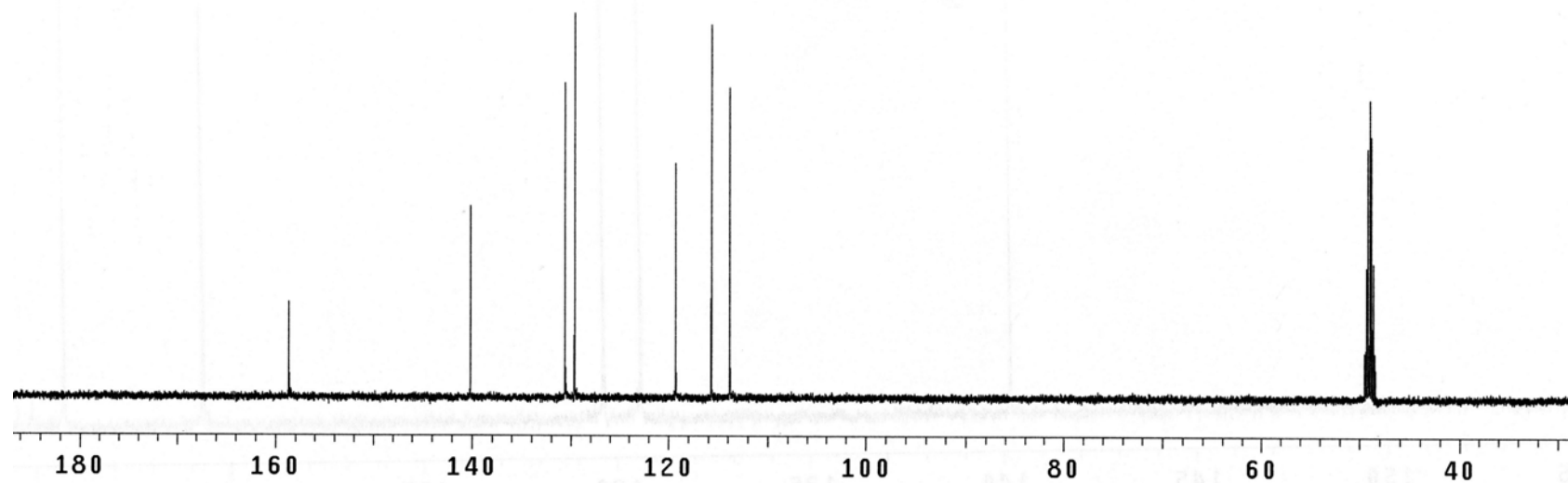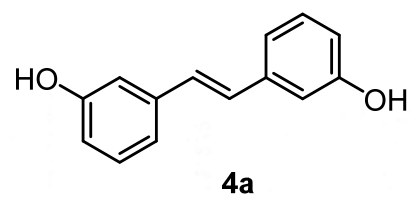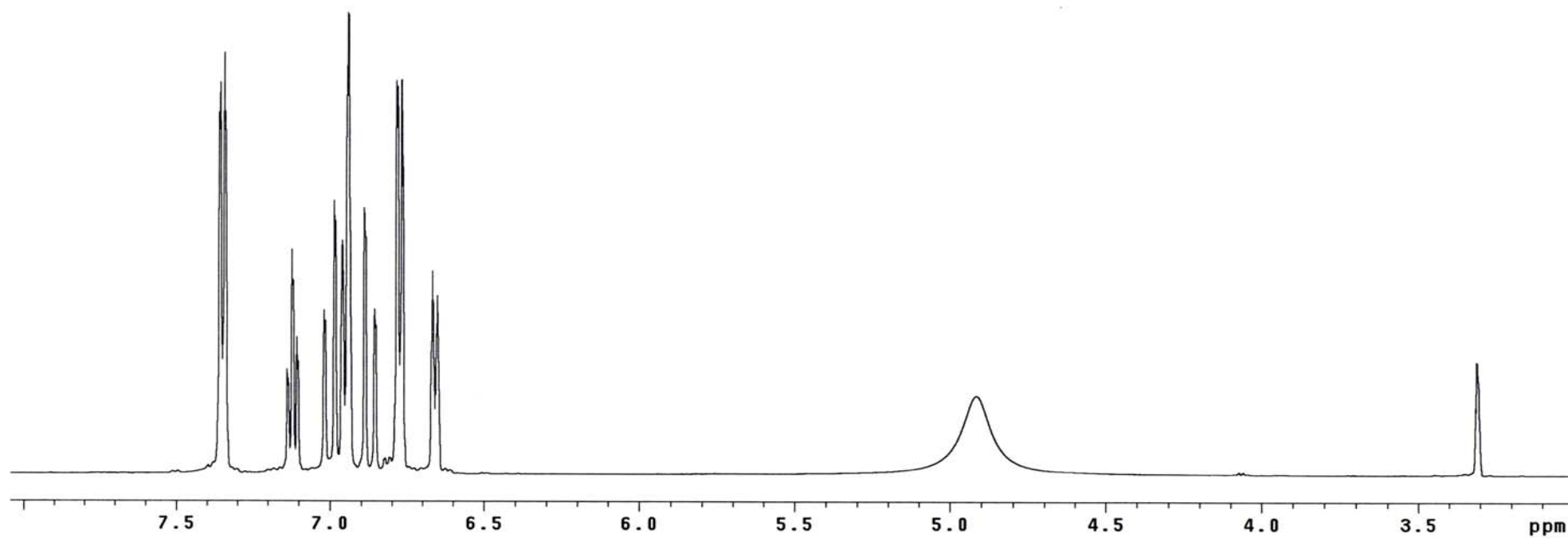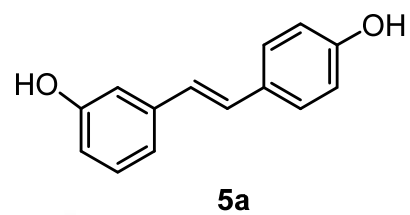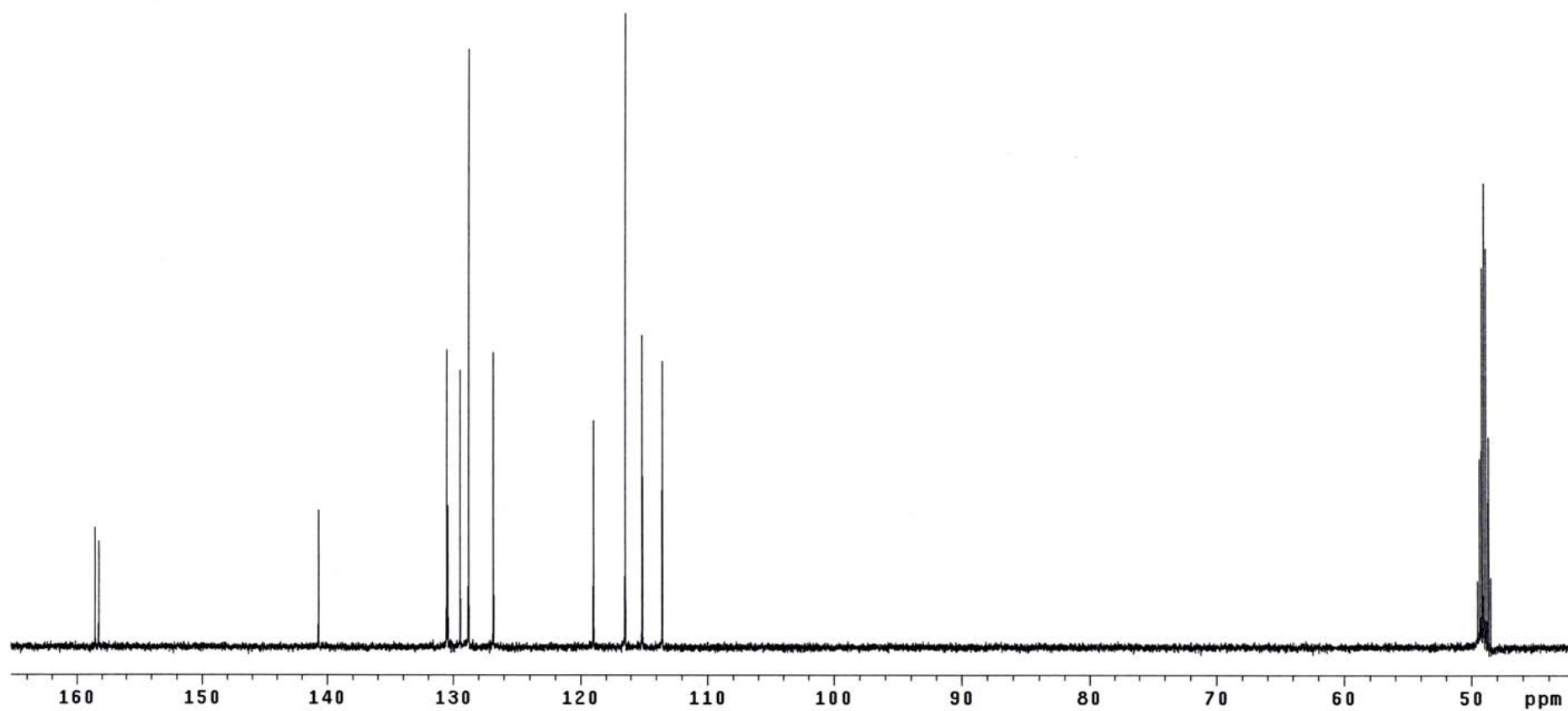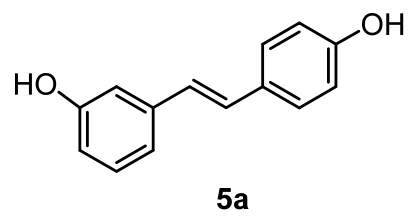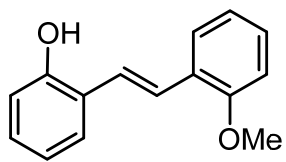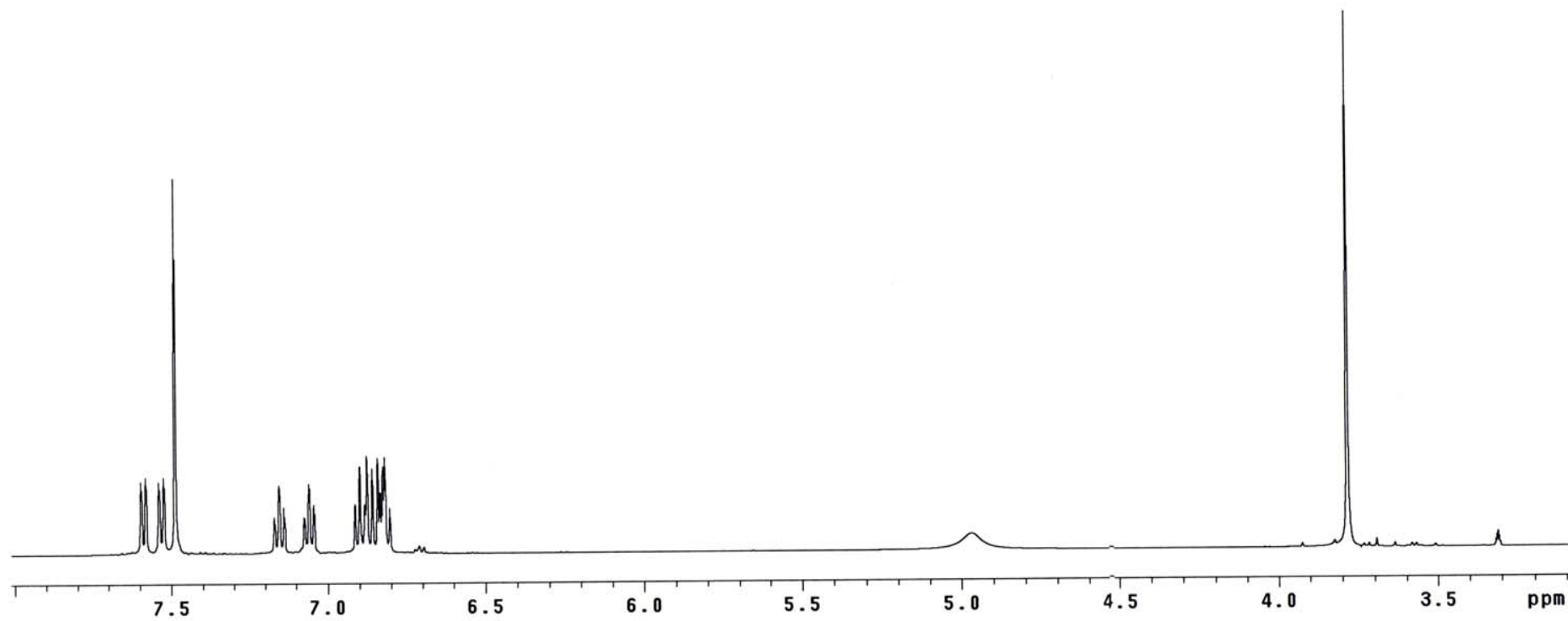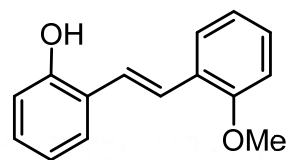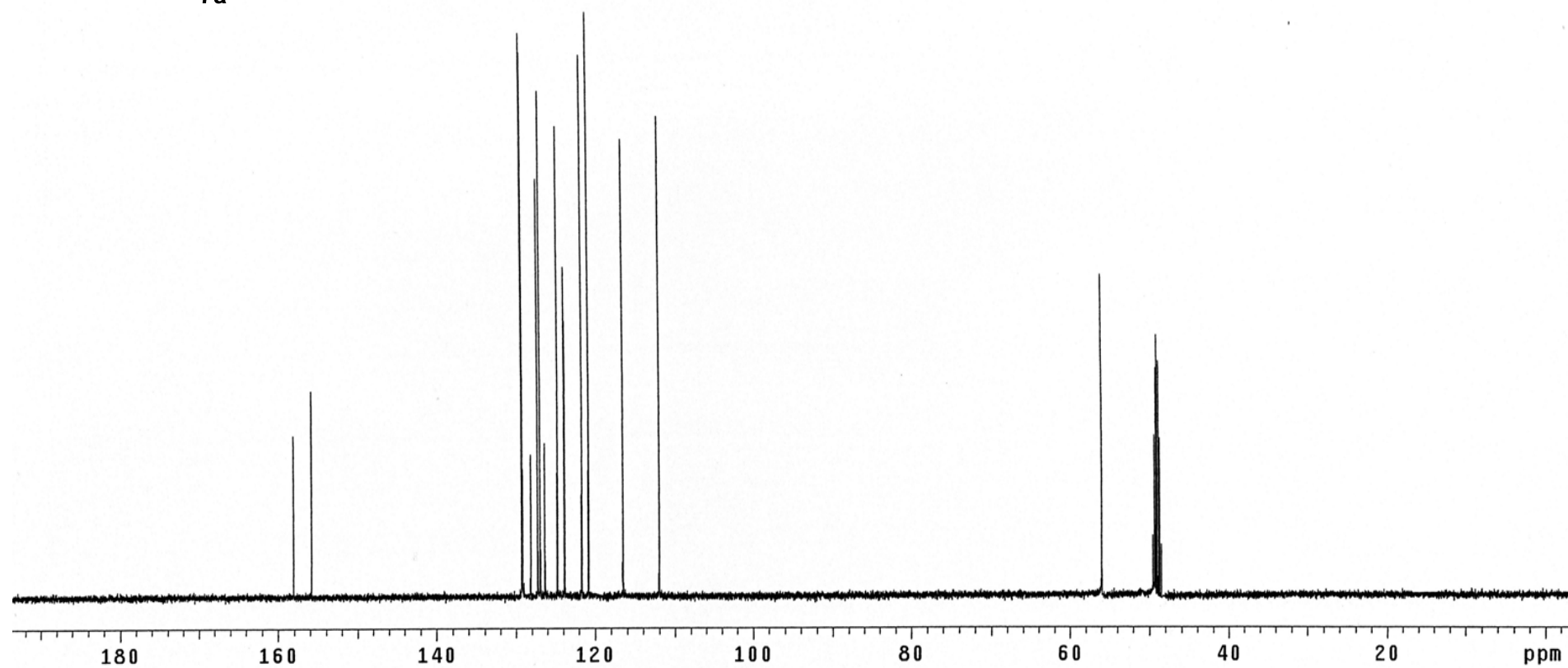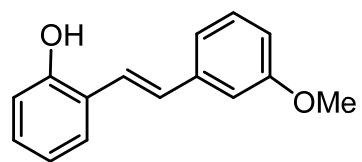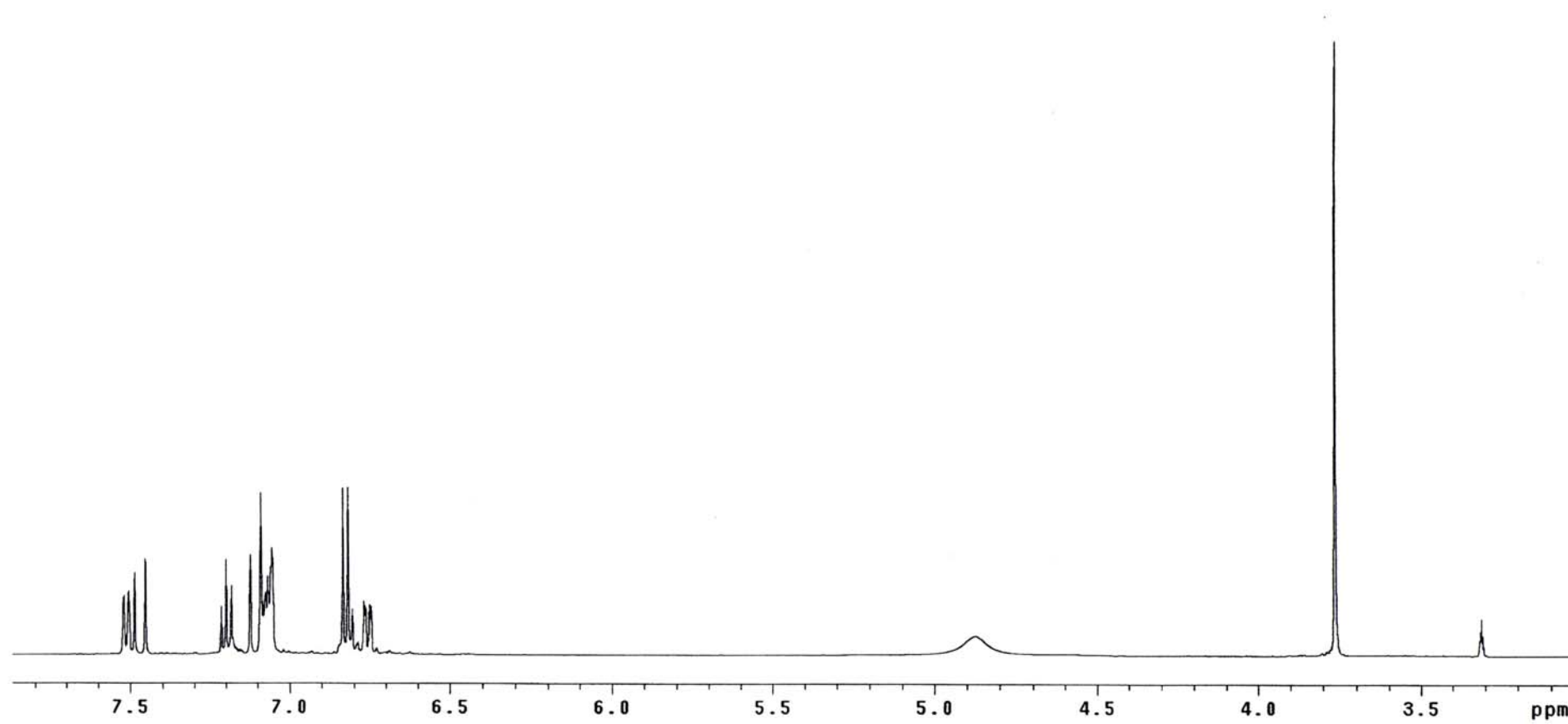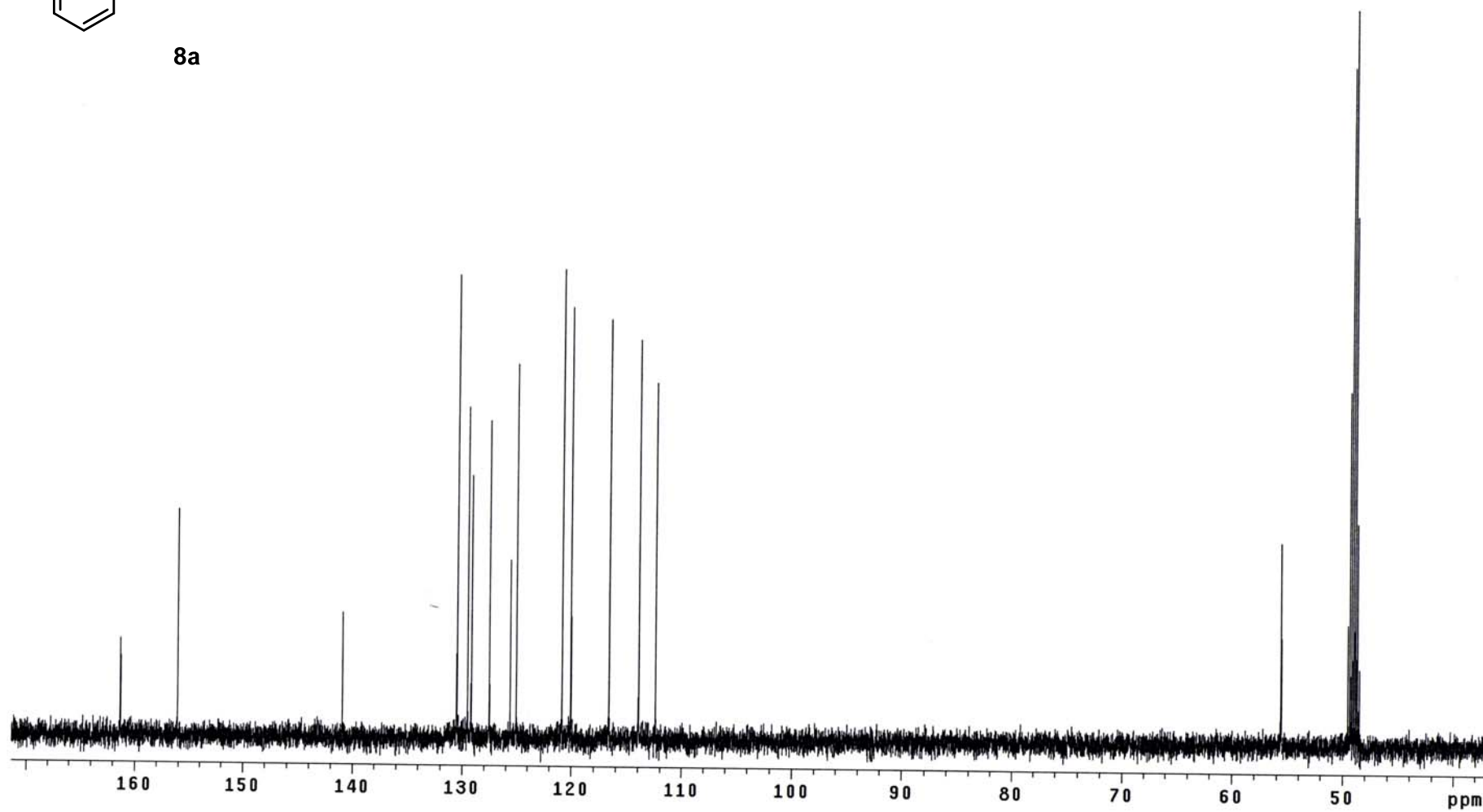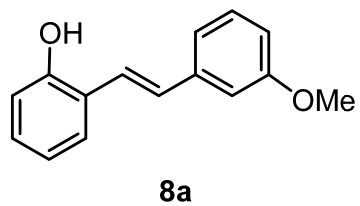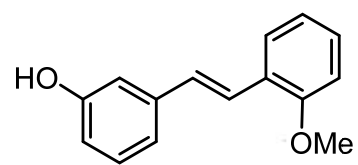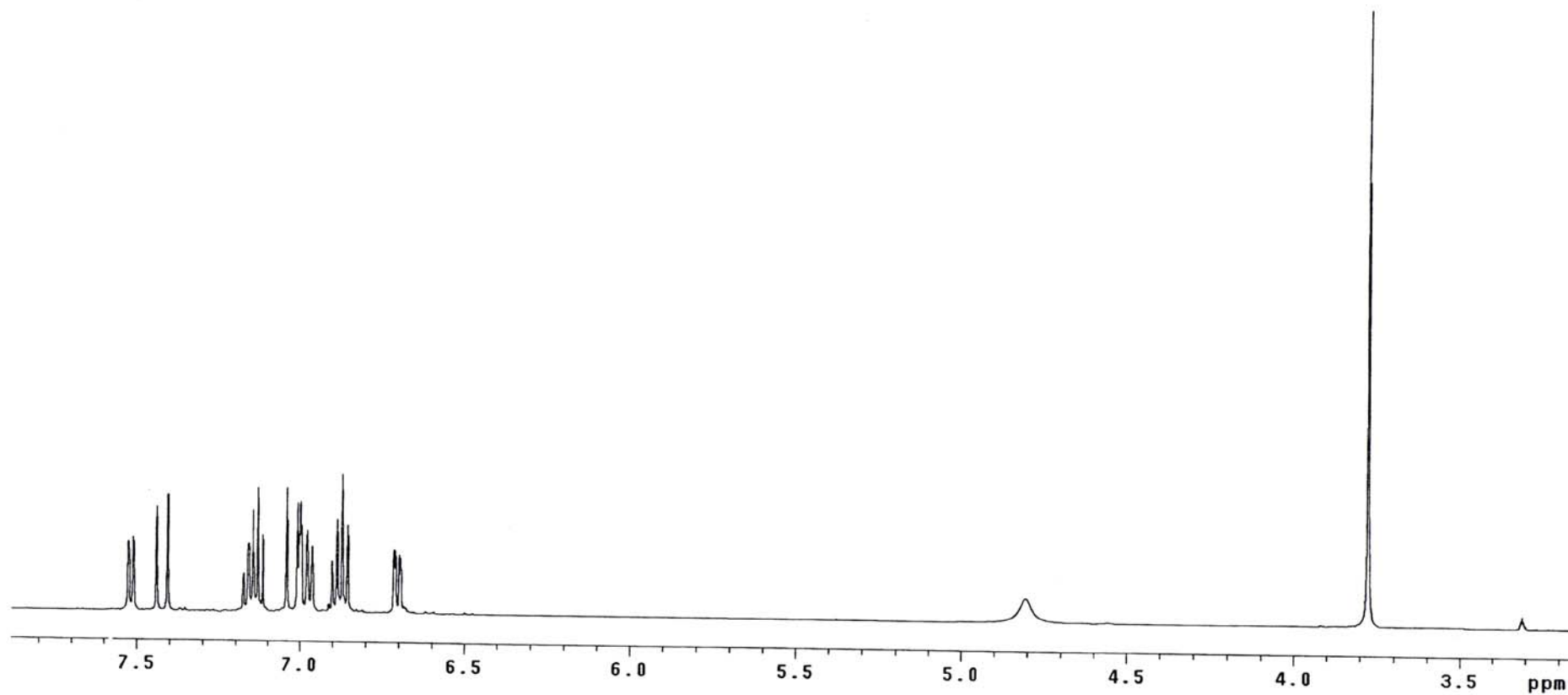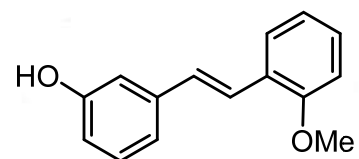
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19b

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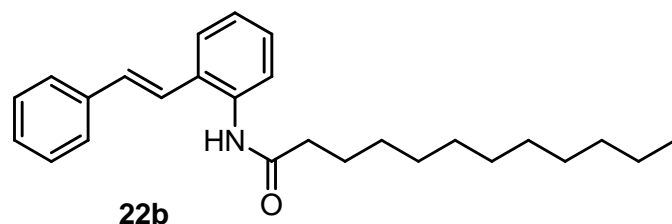160   150   140   130   120   110   100   90   80   70   60   ppm
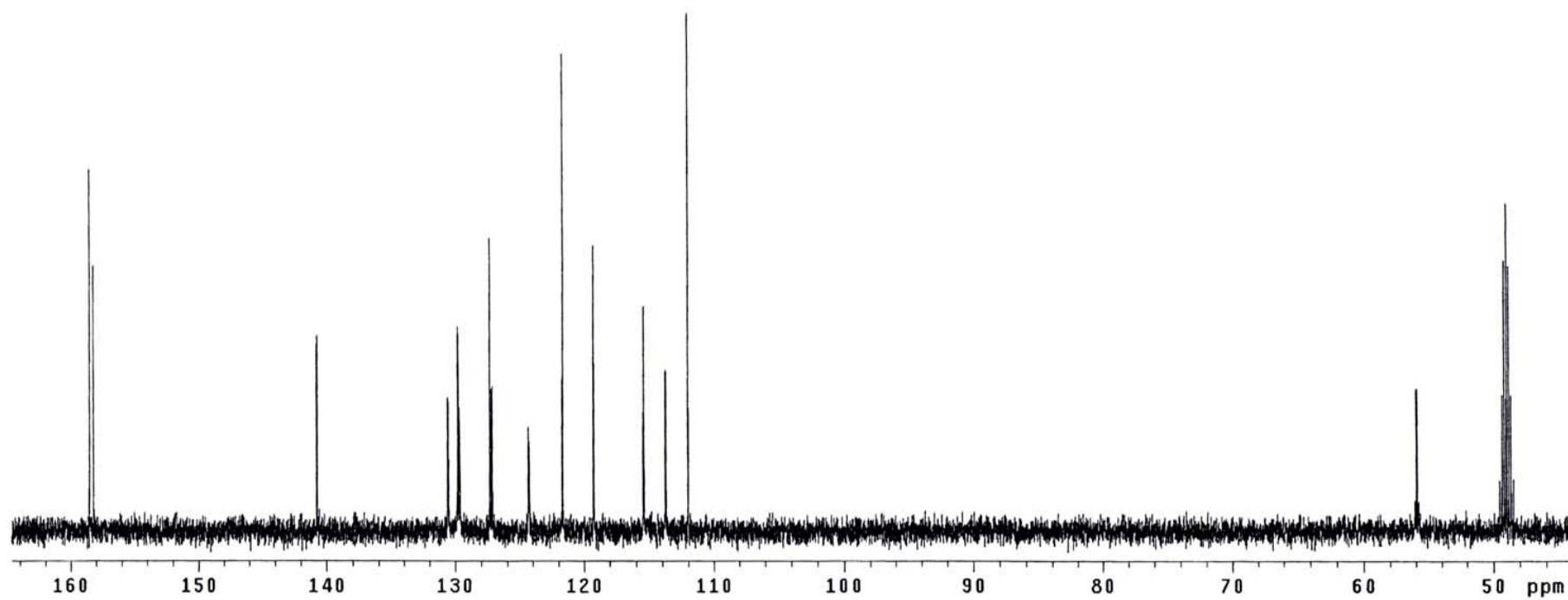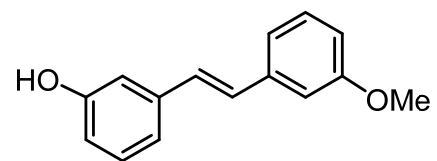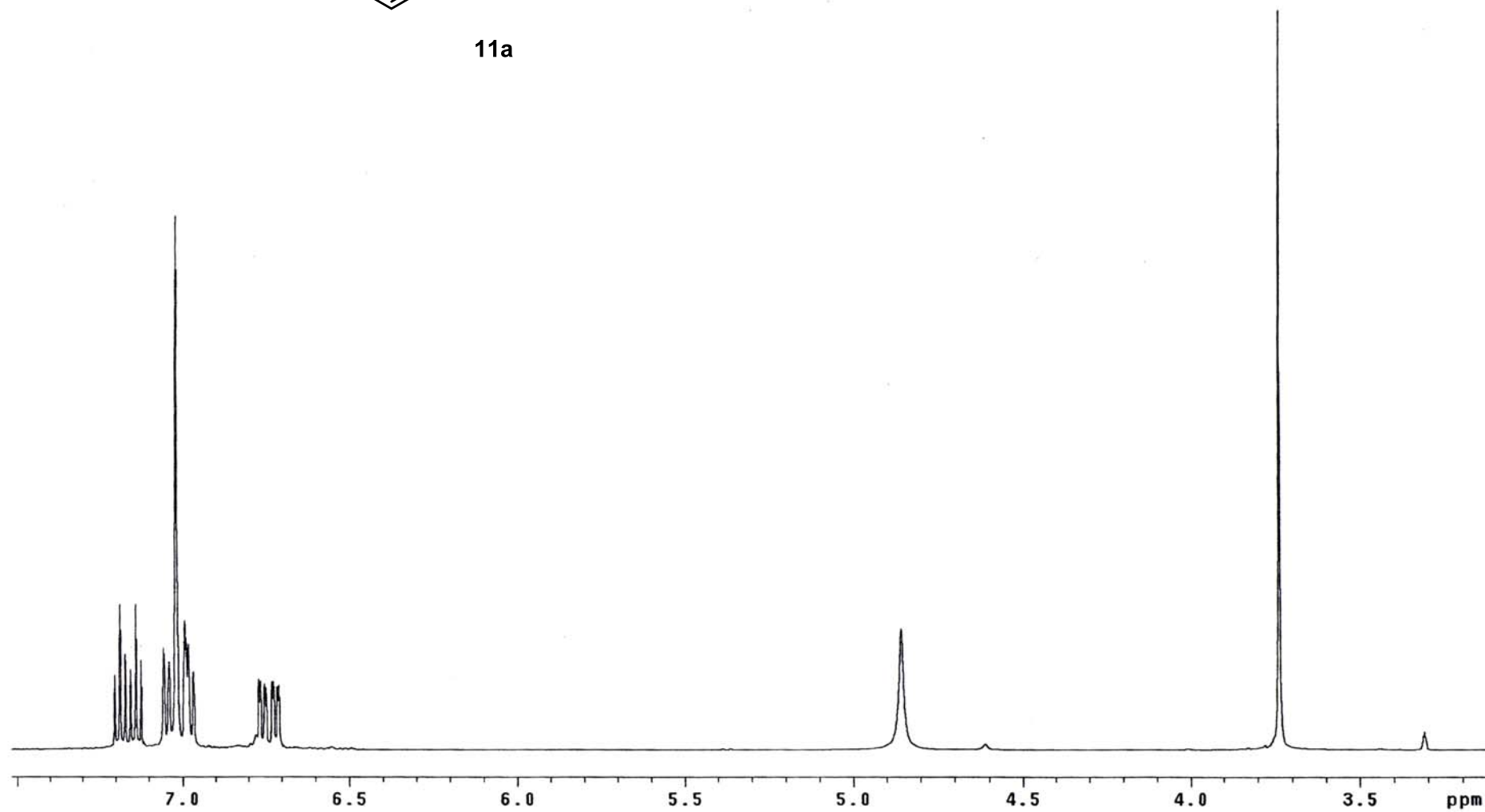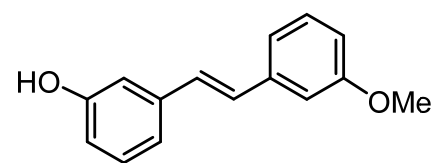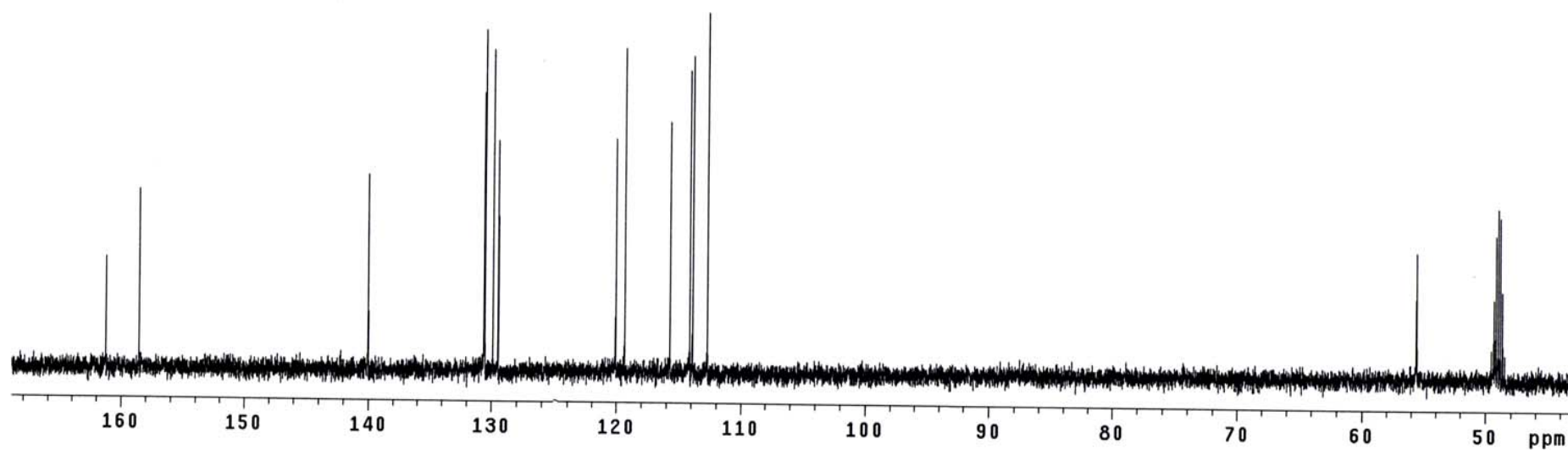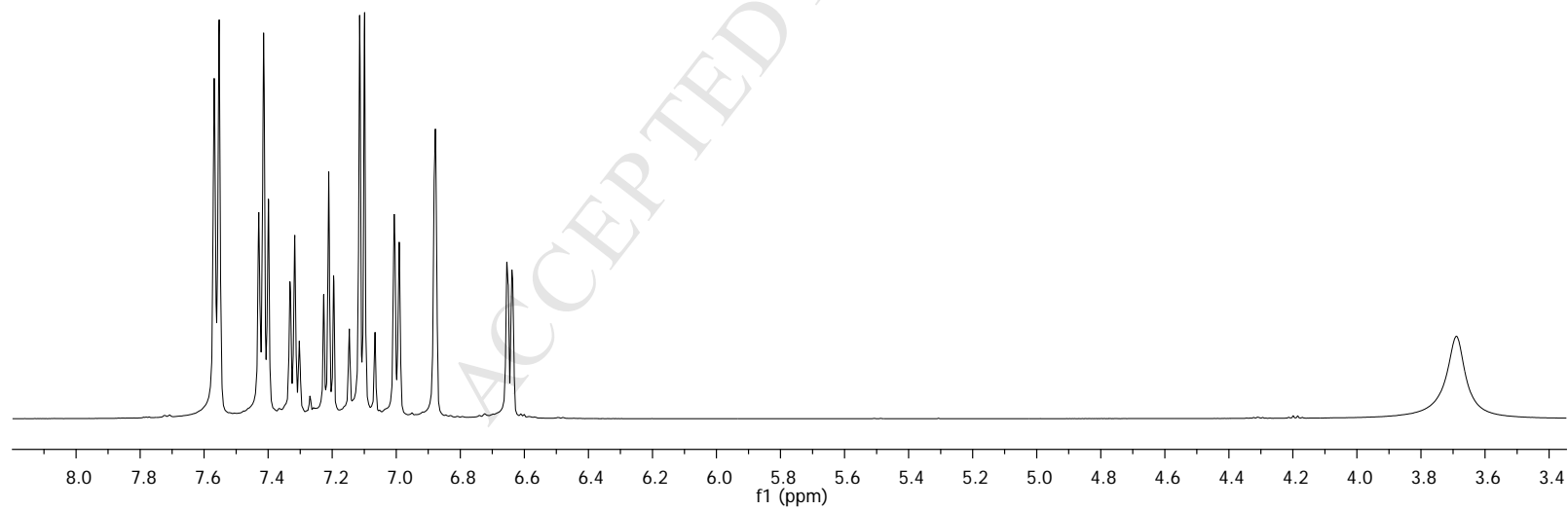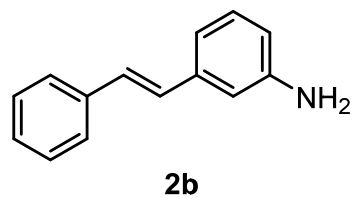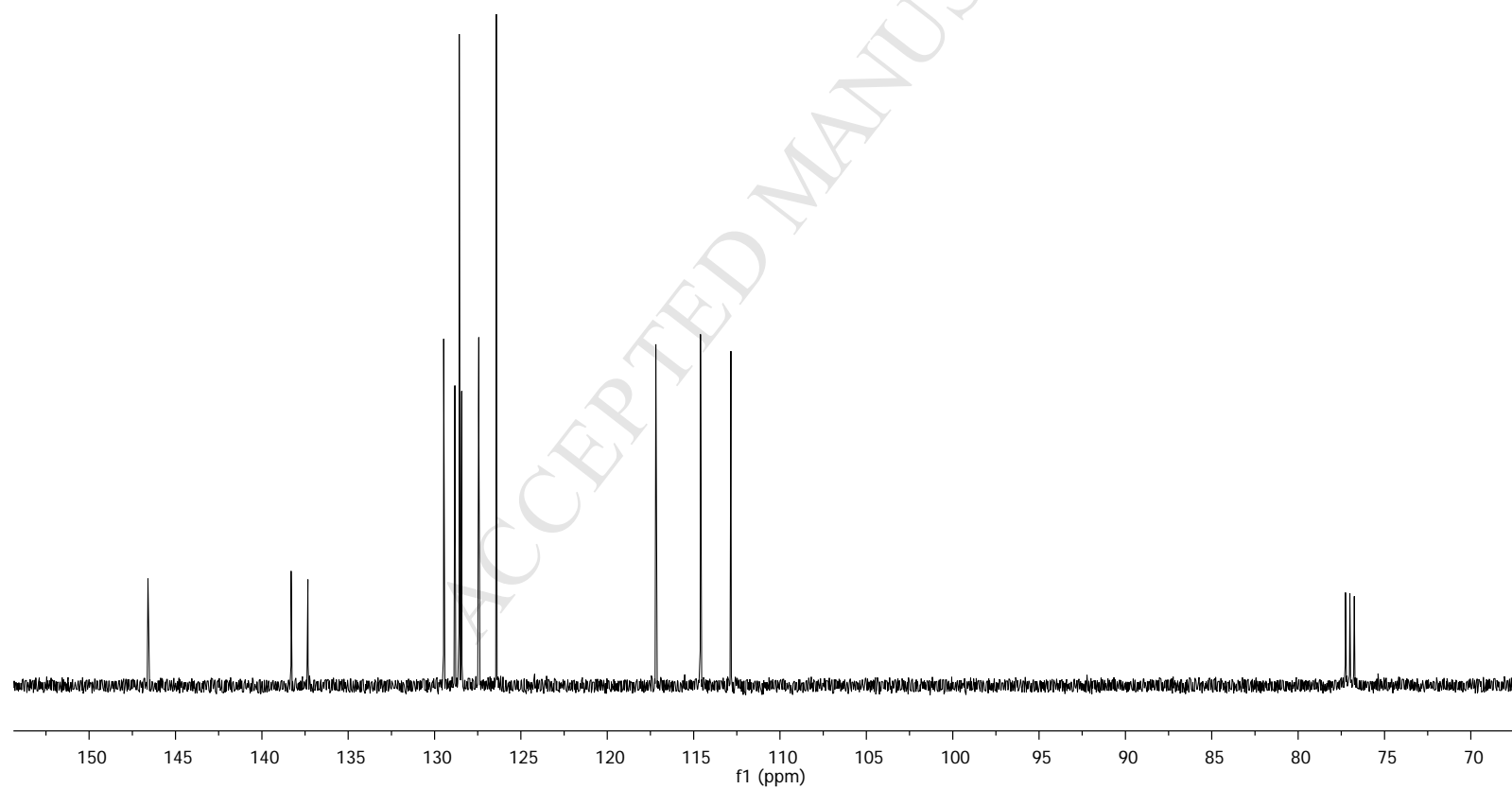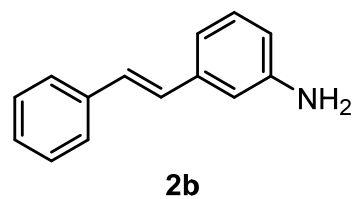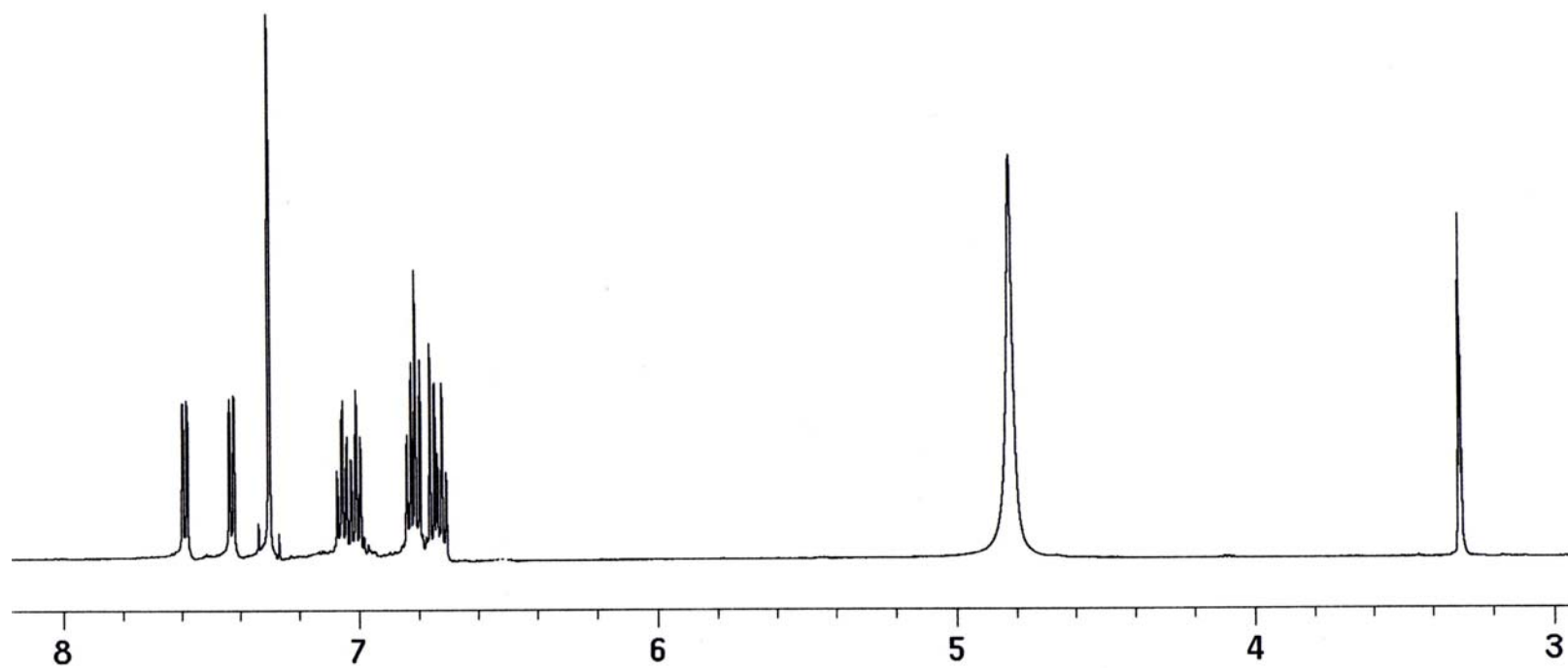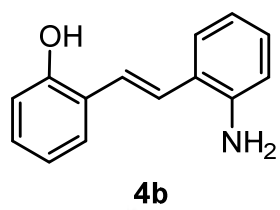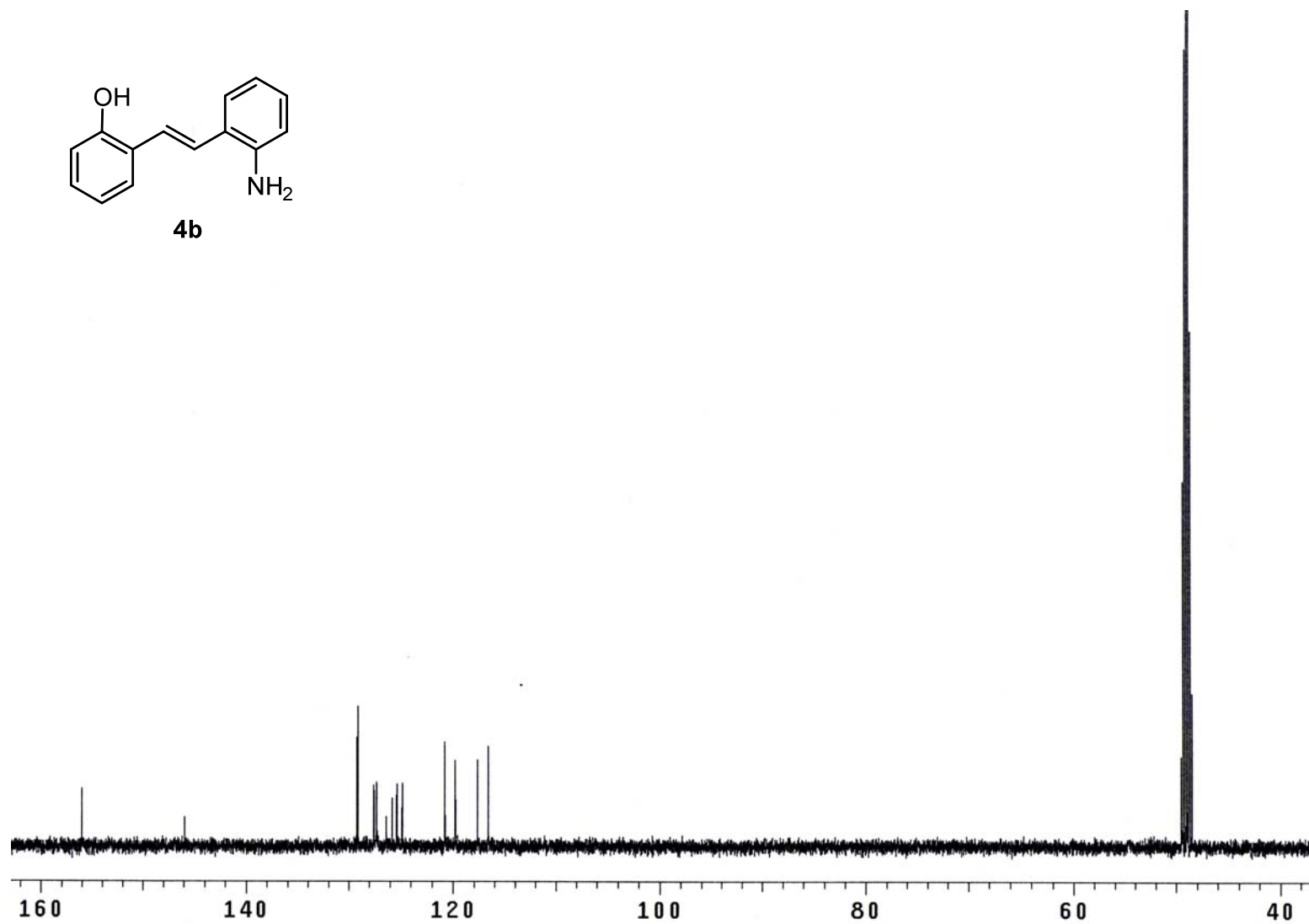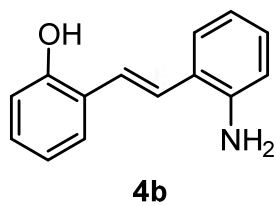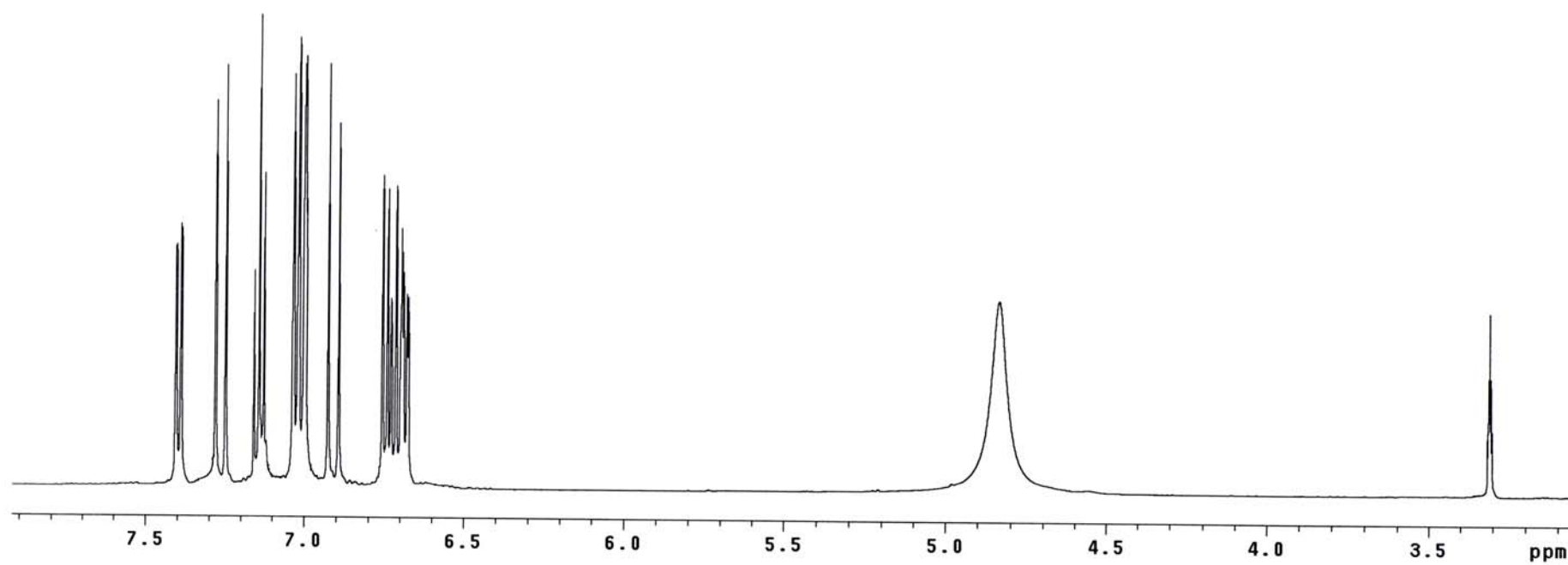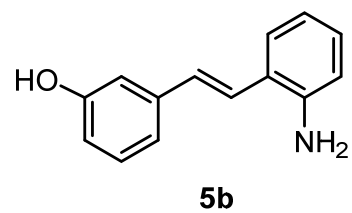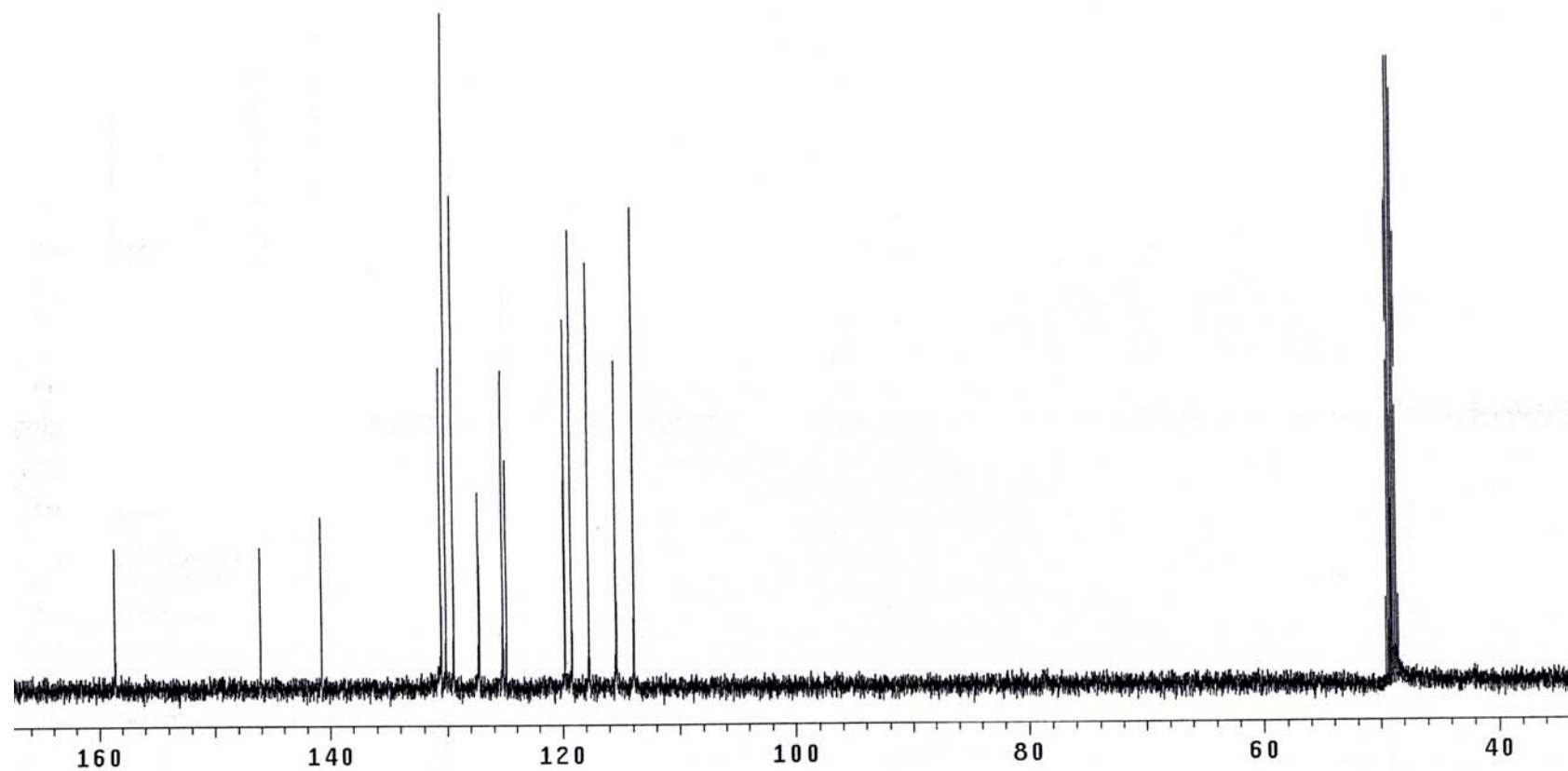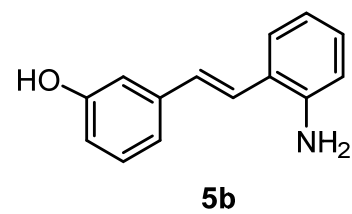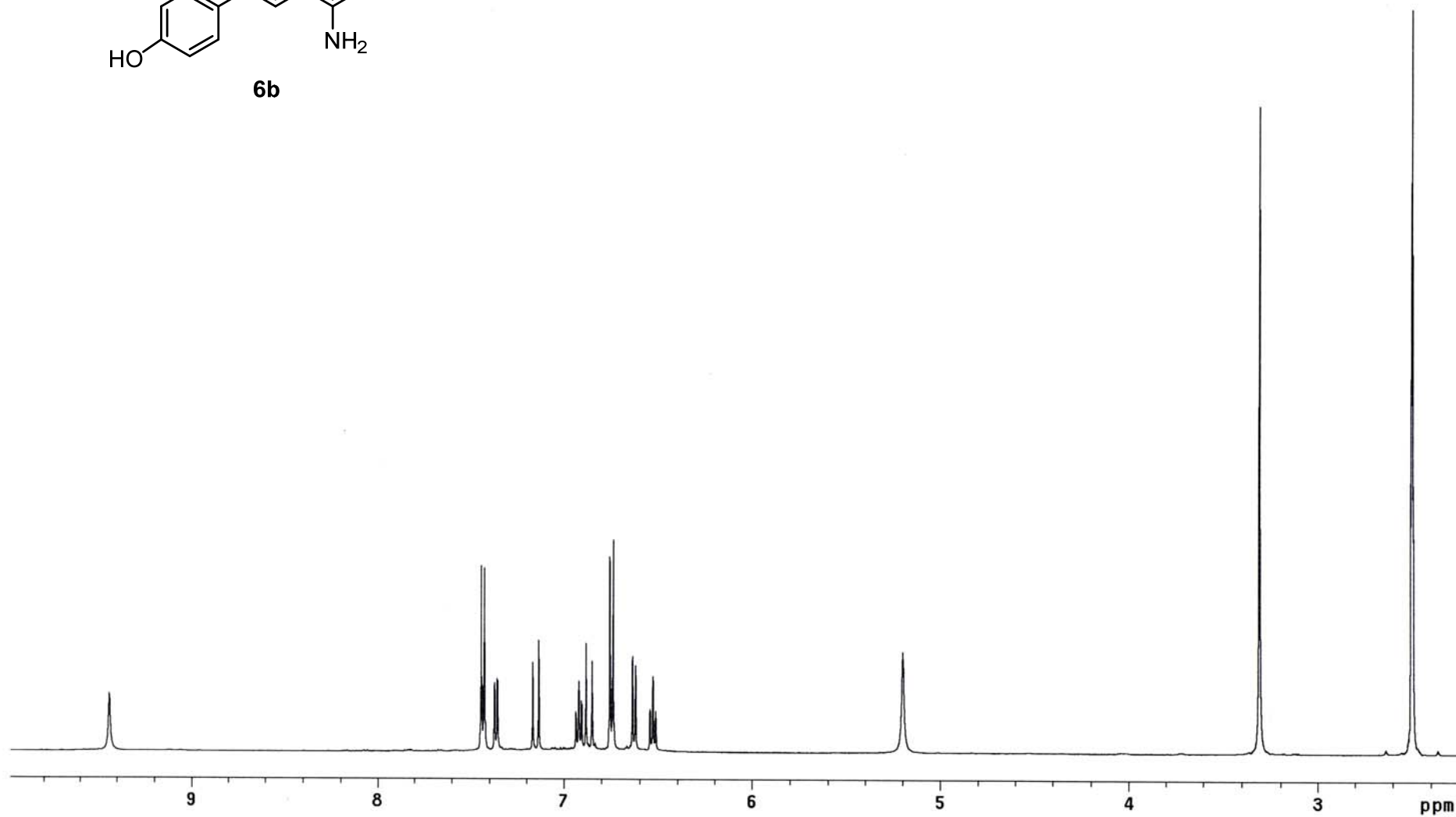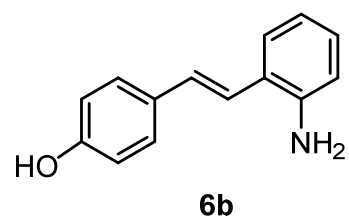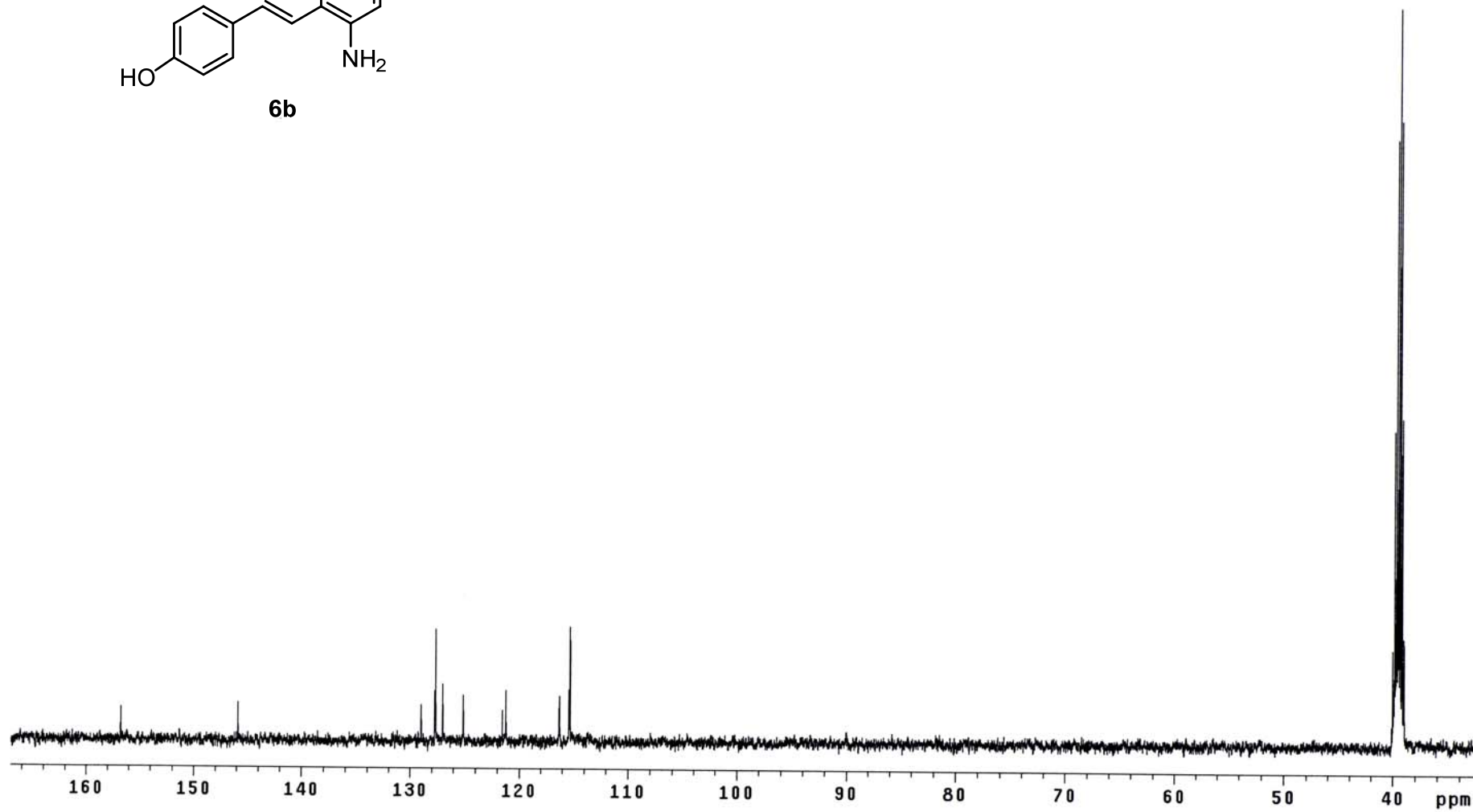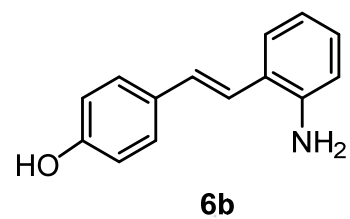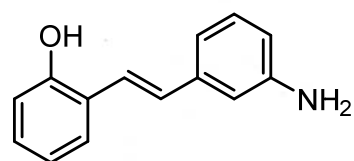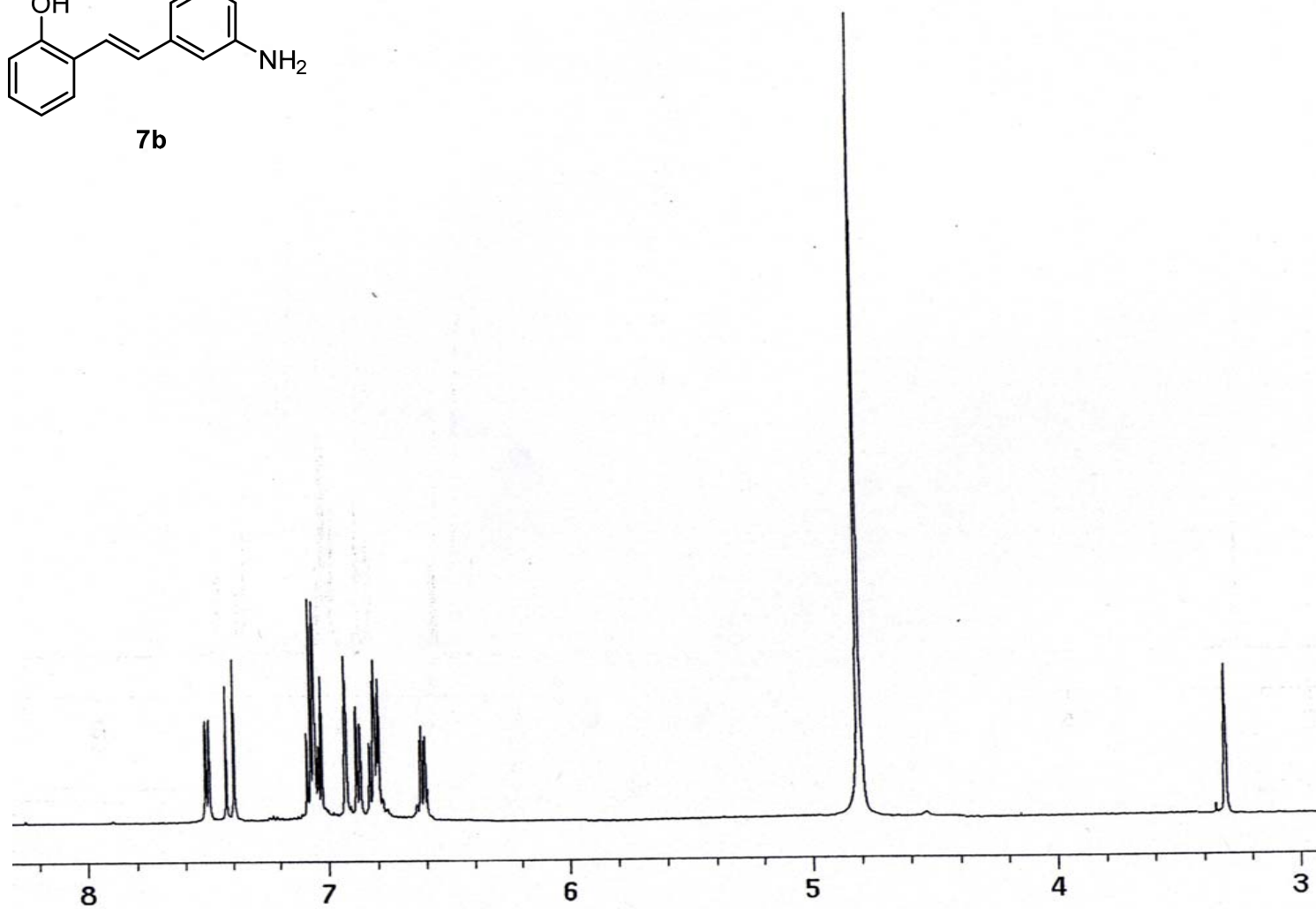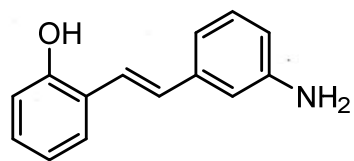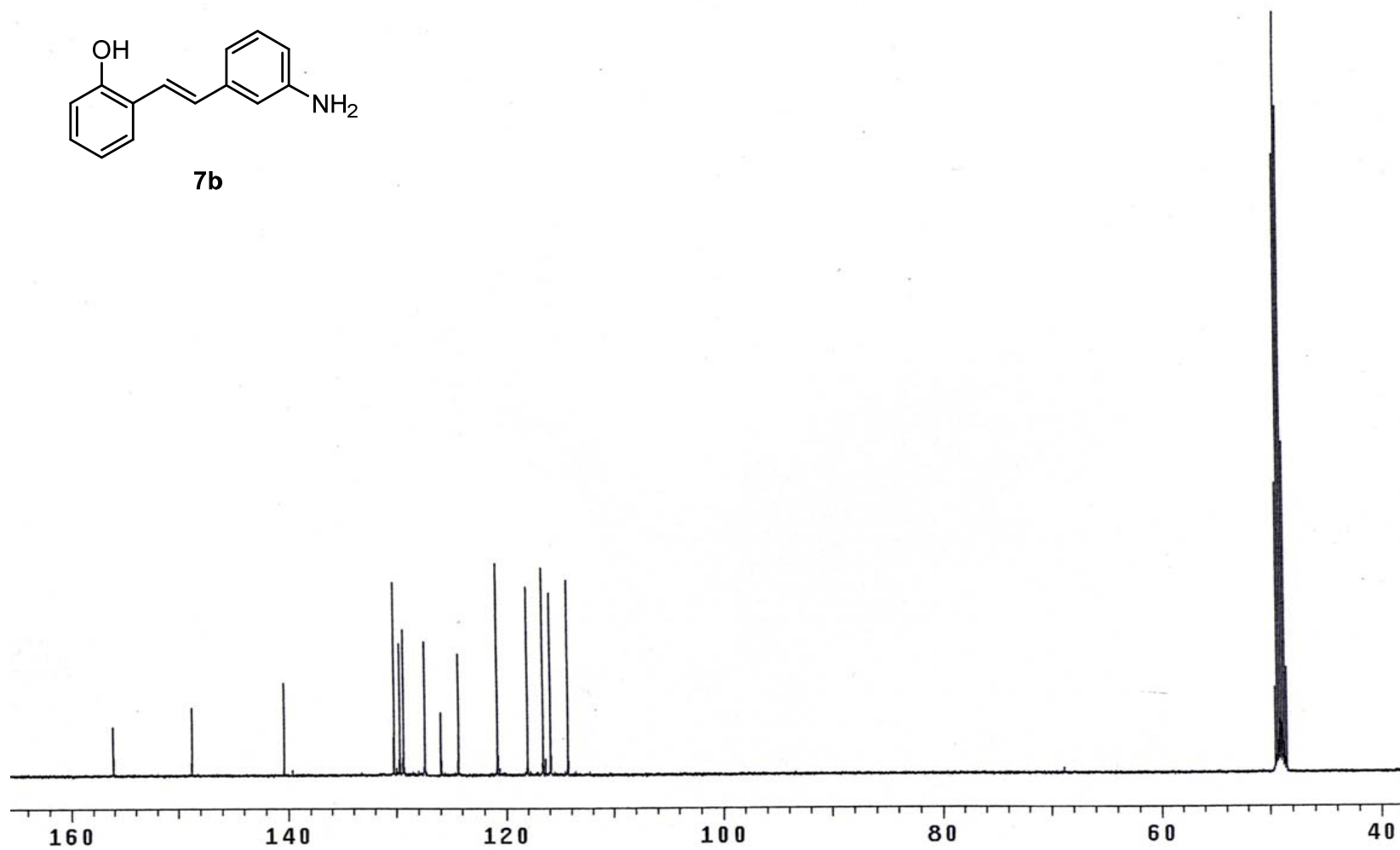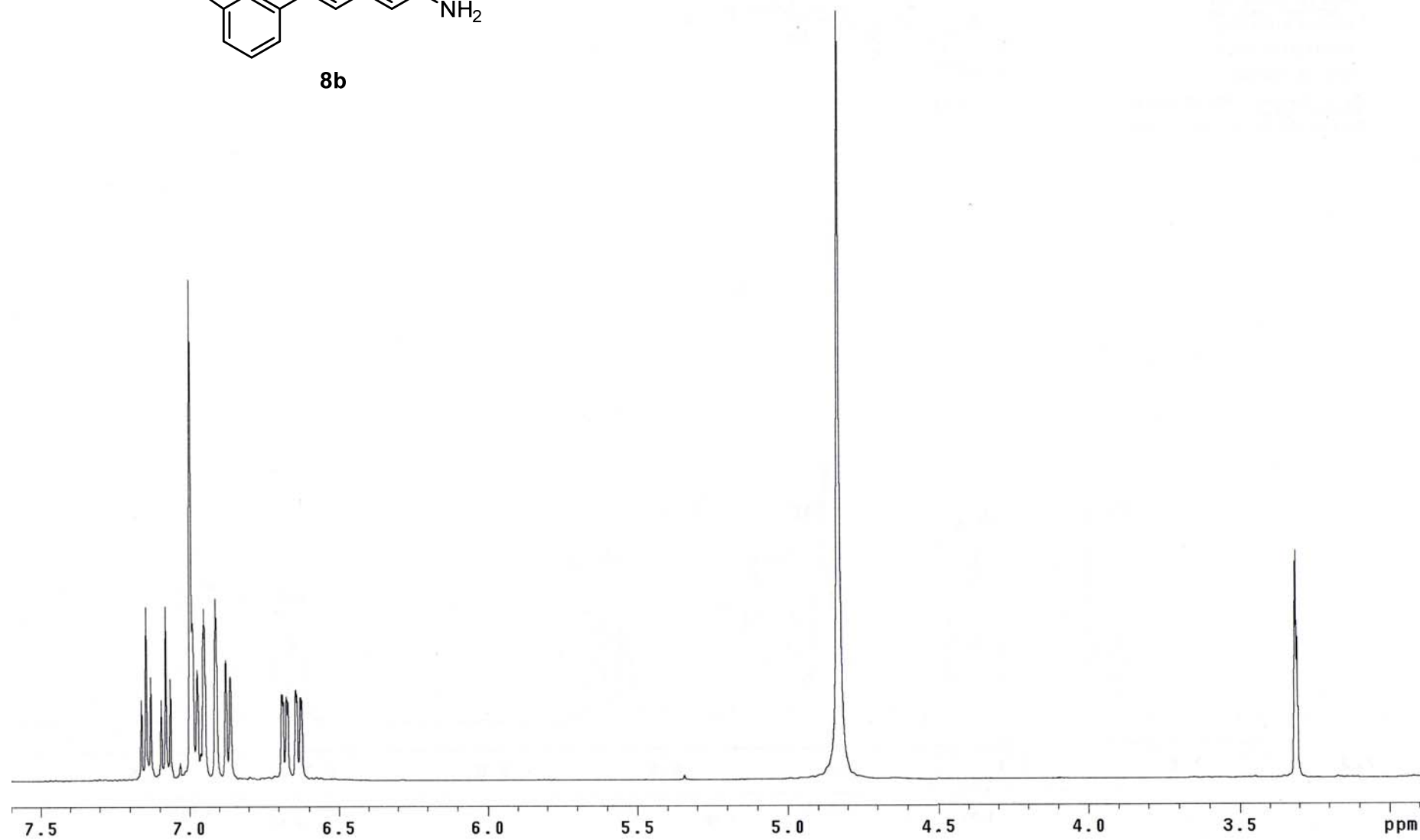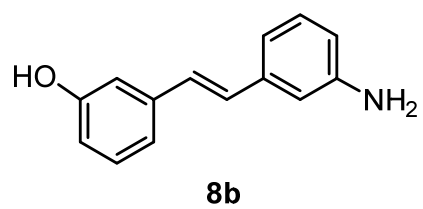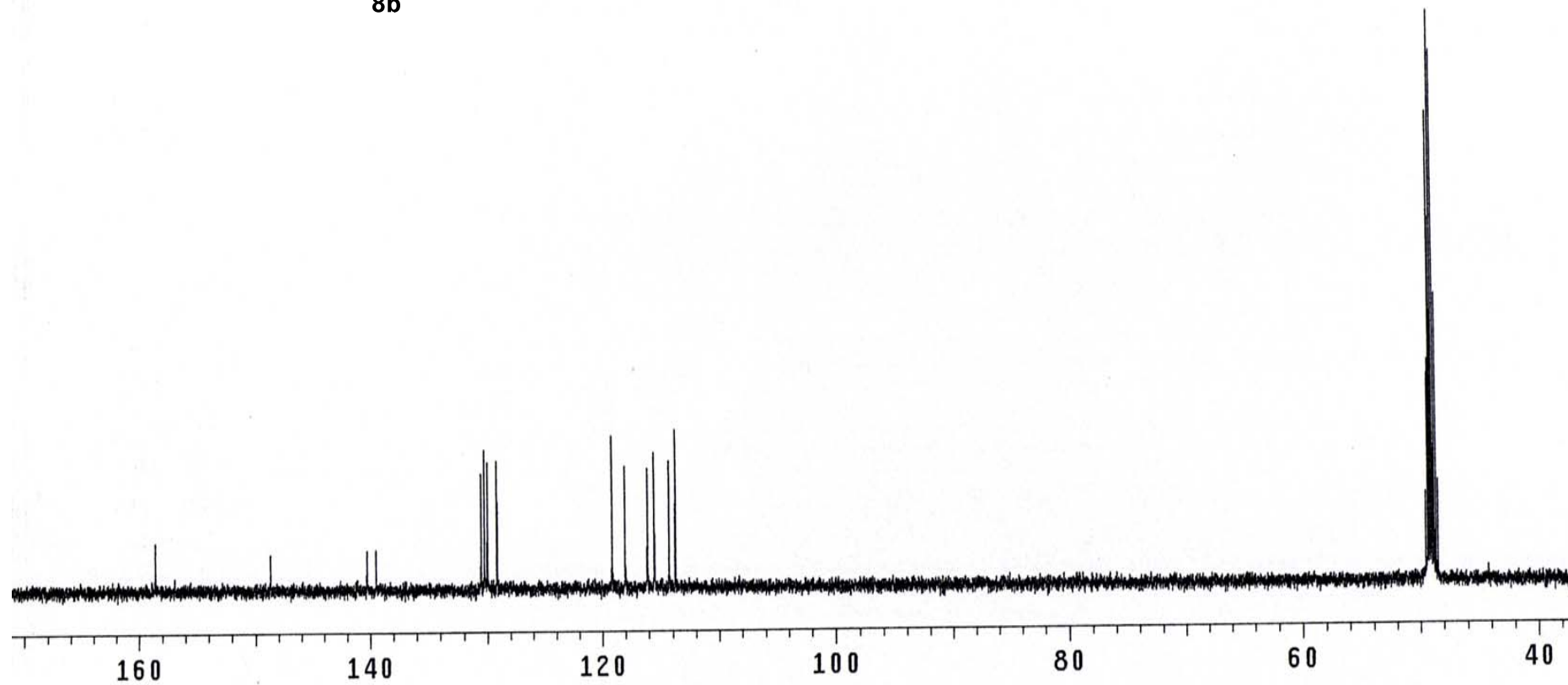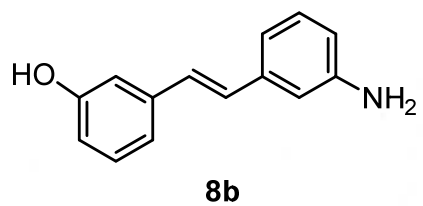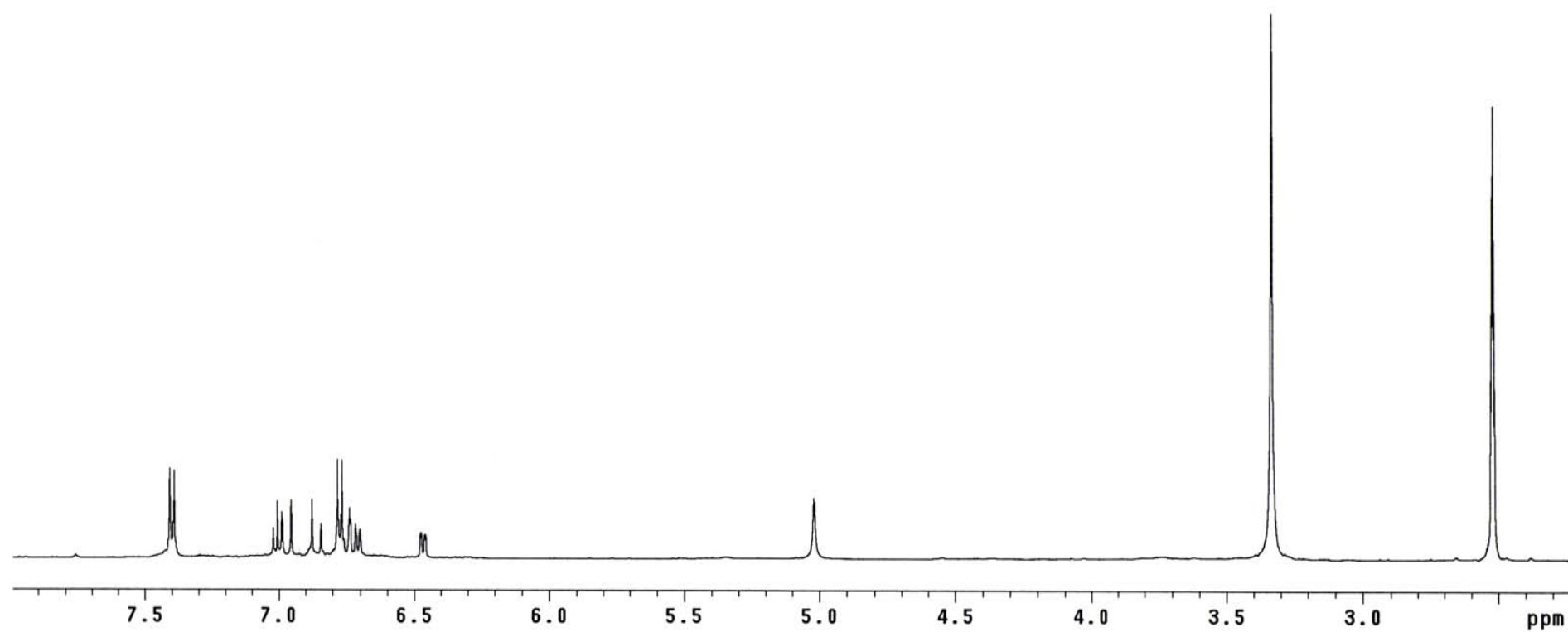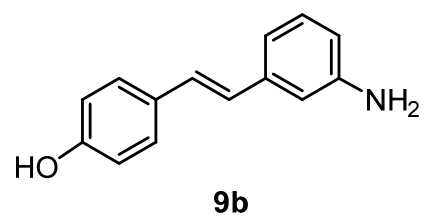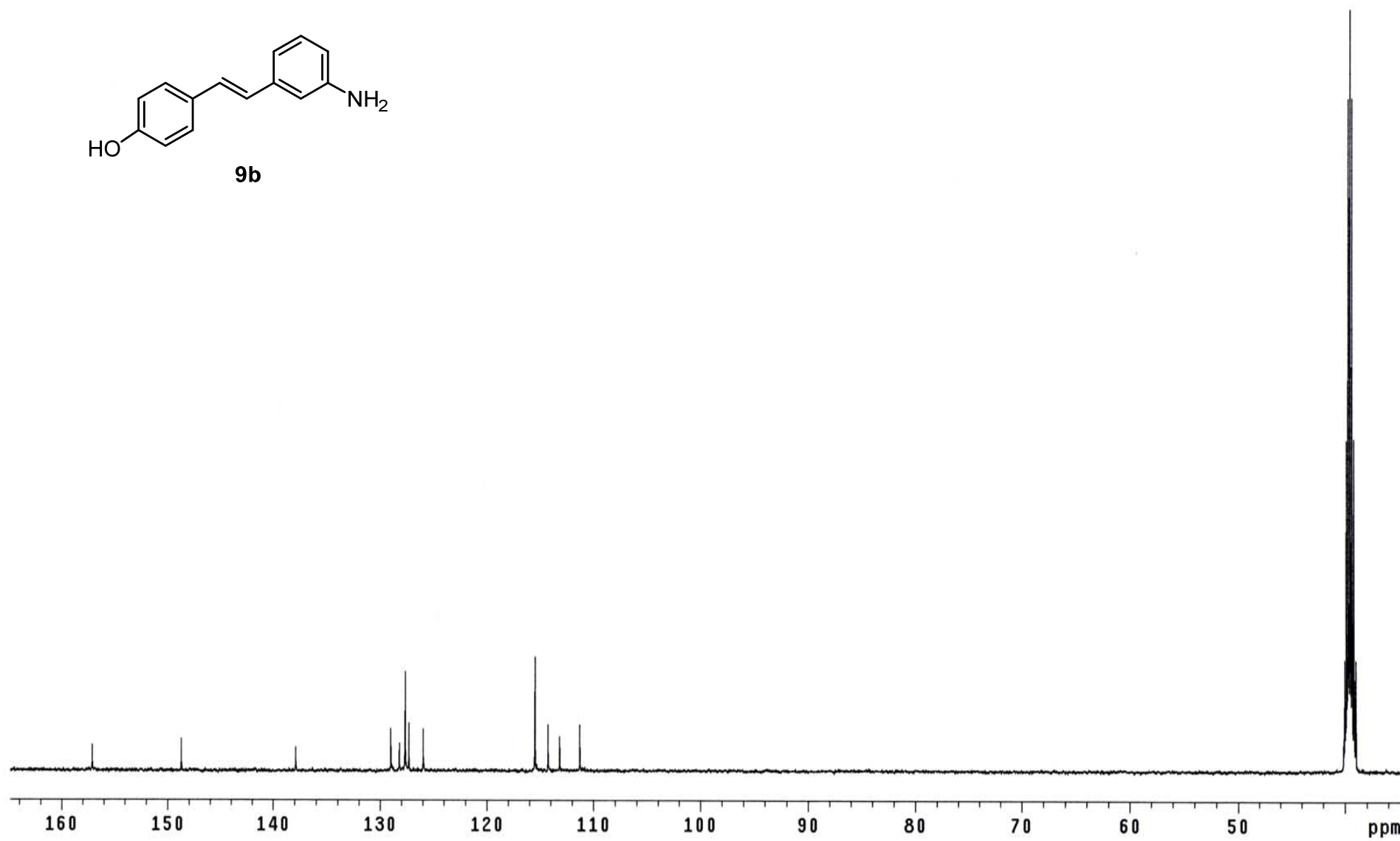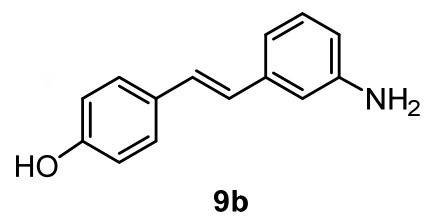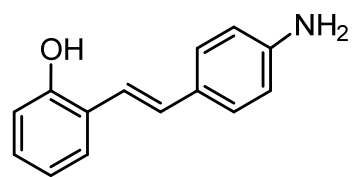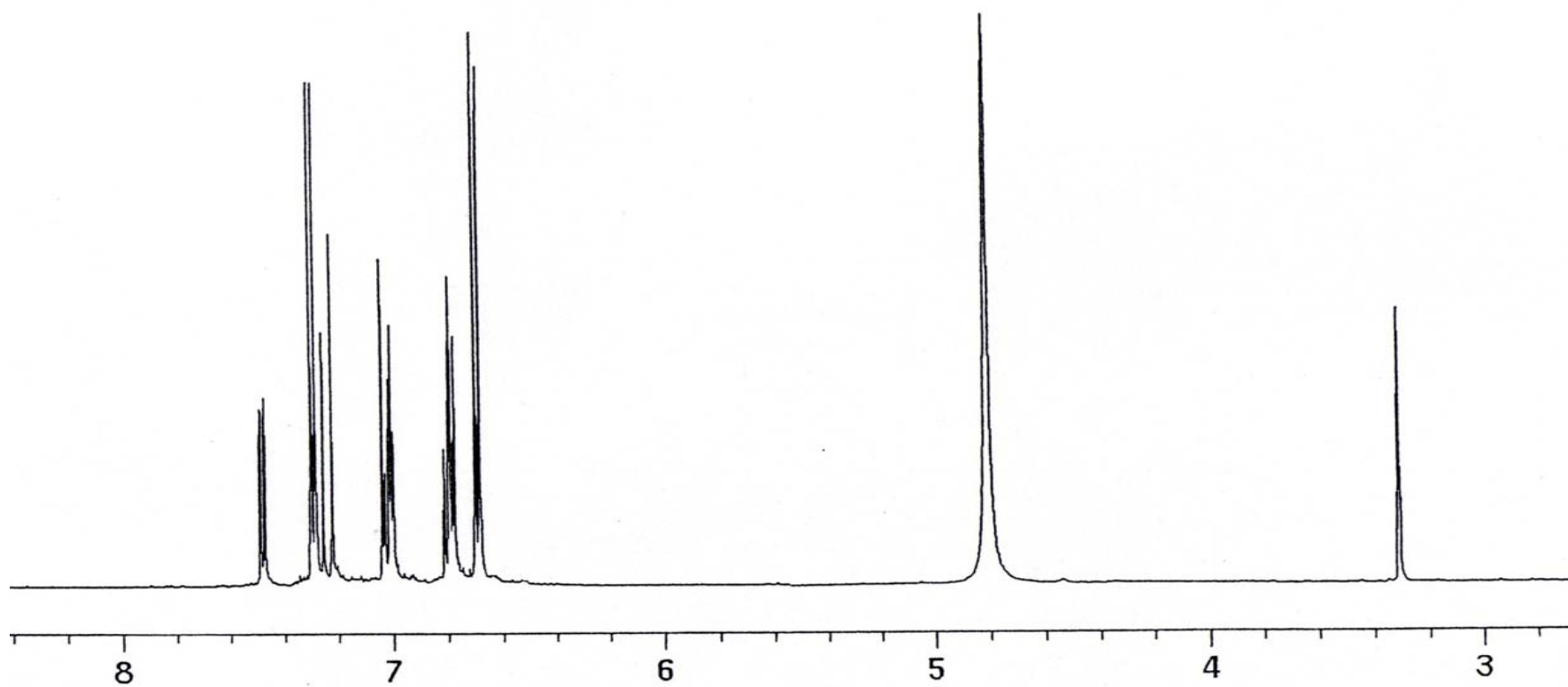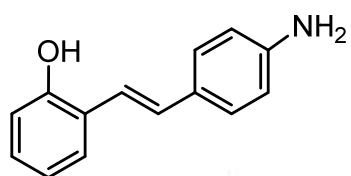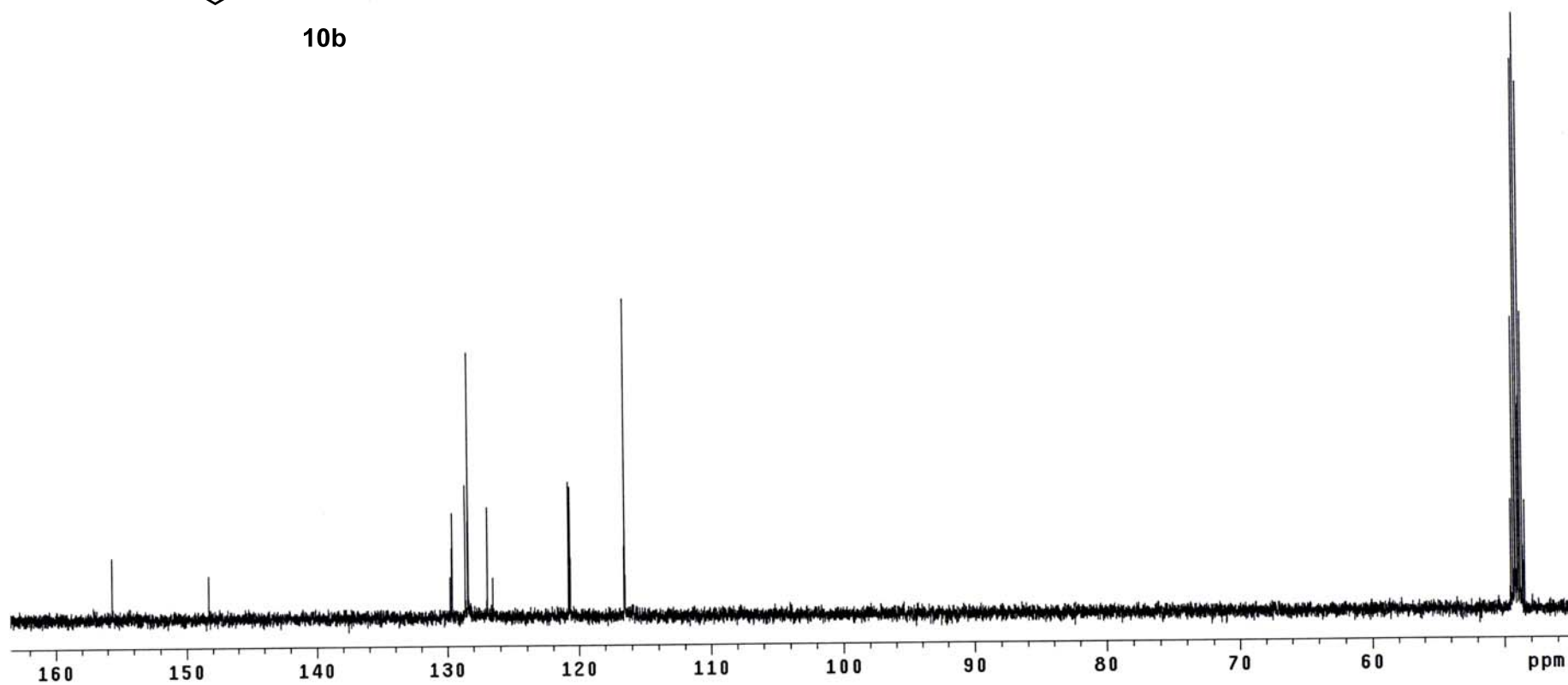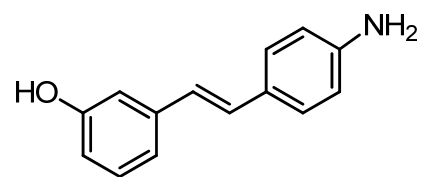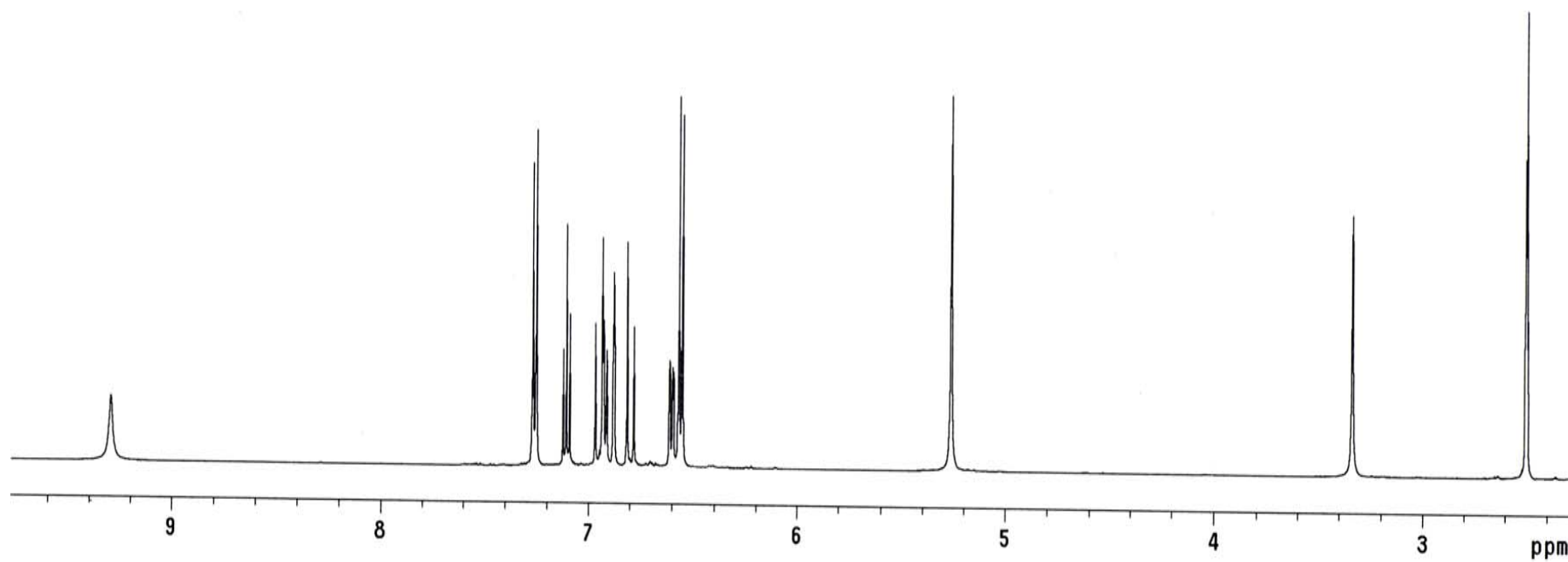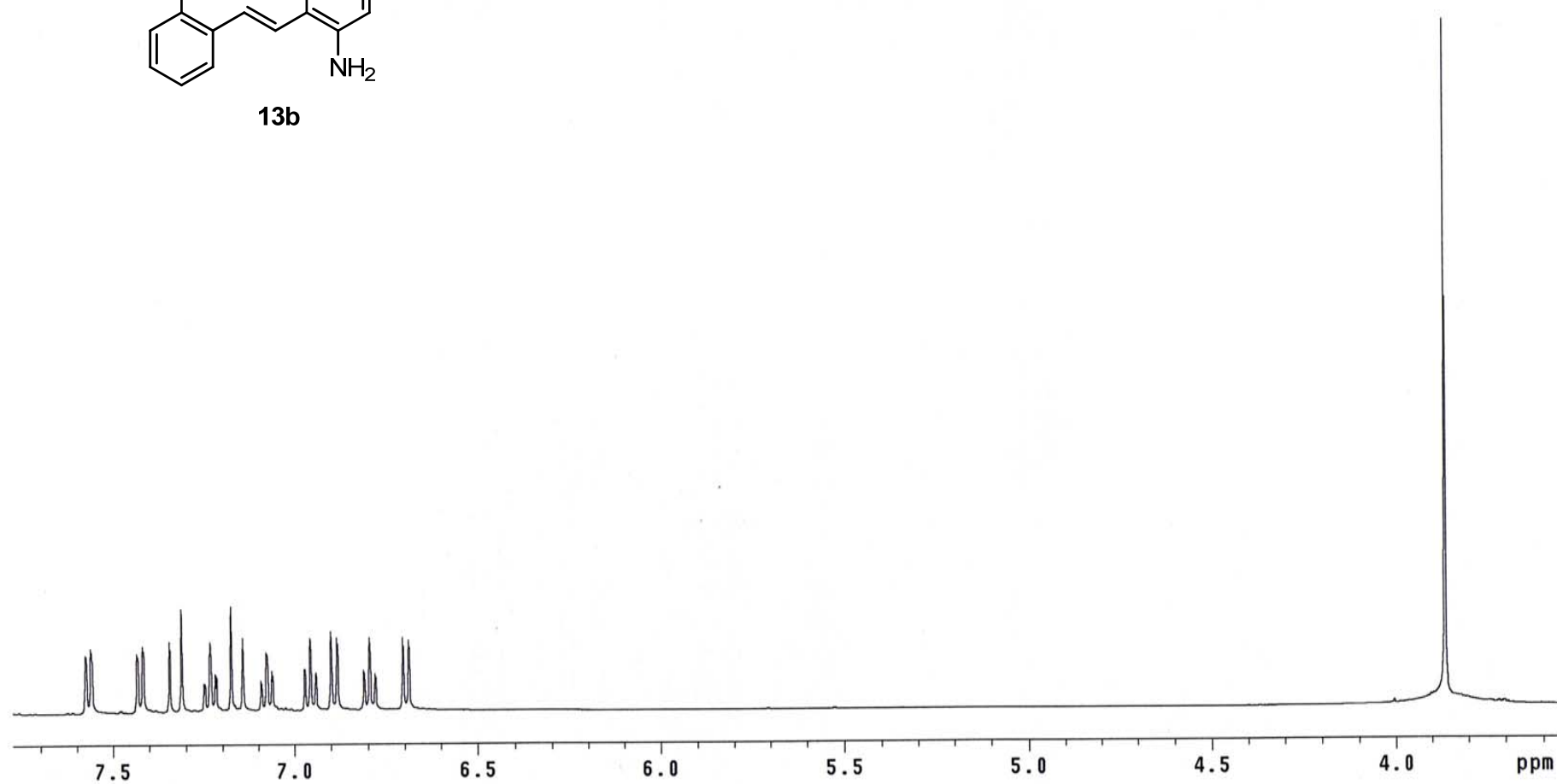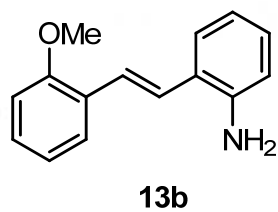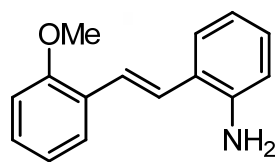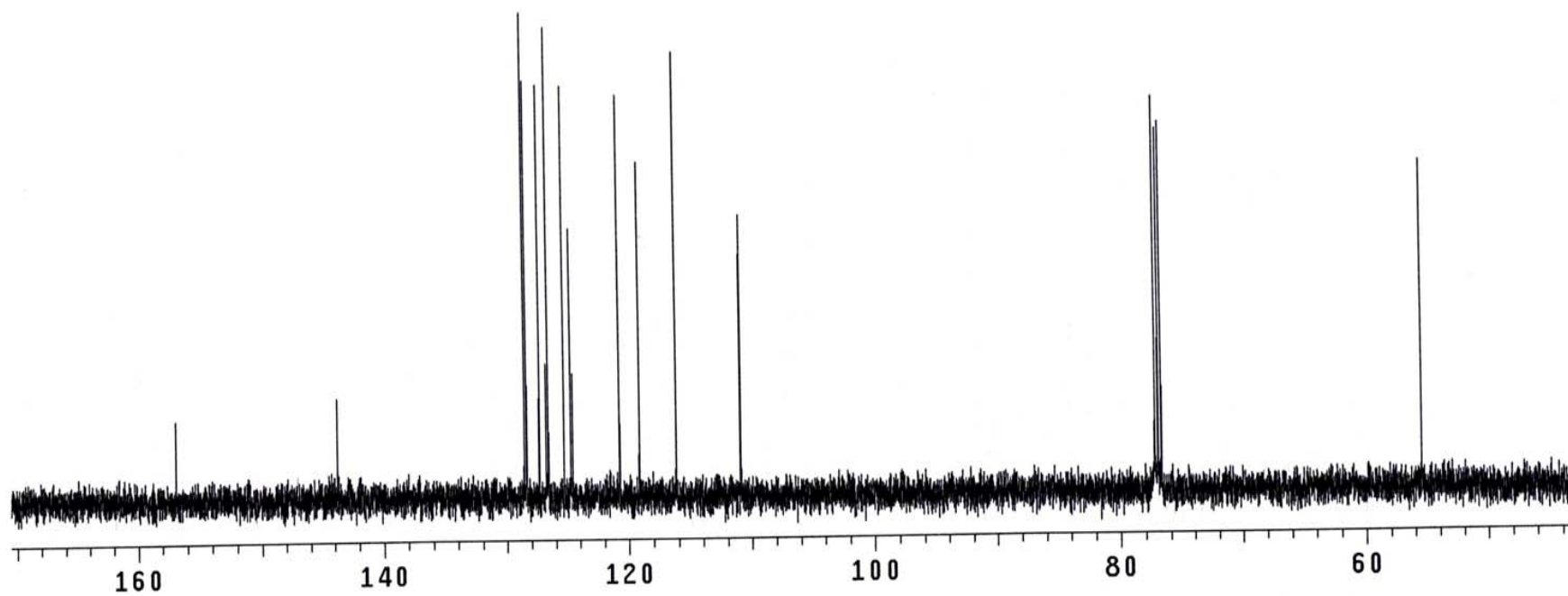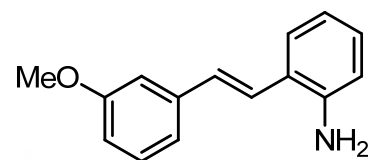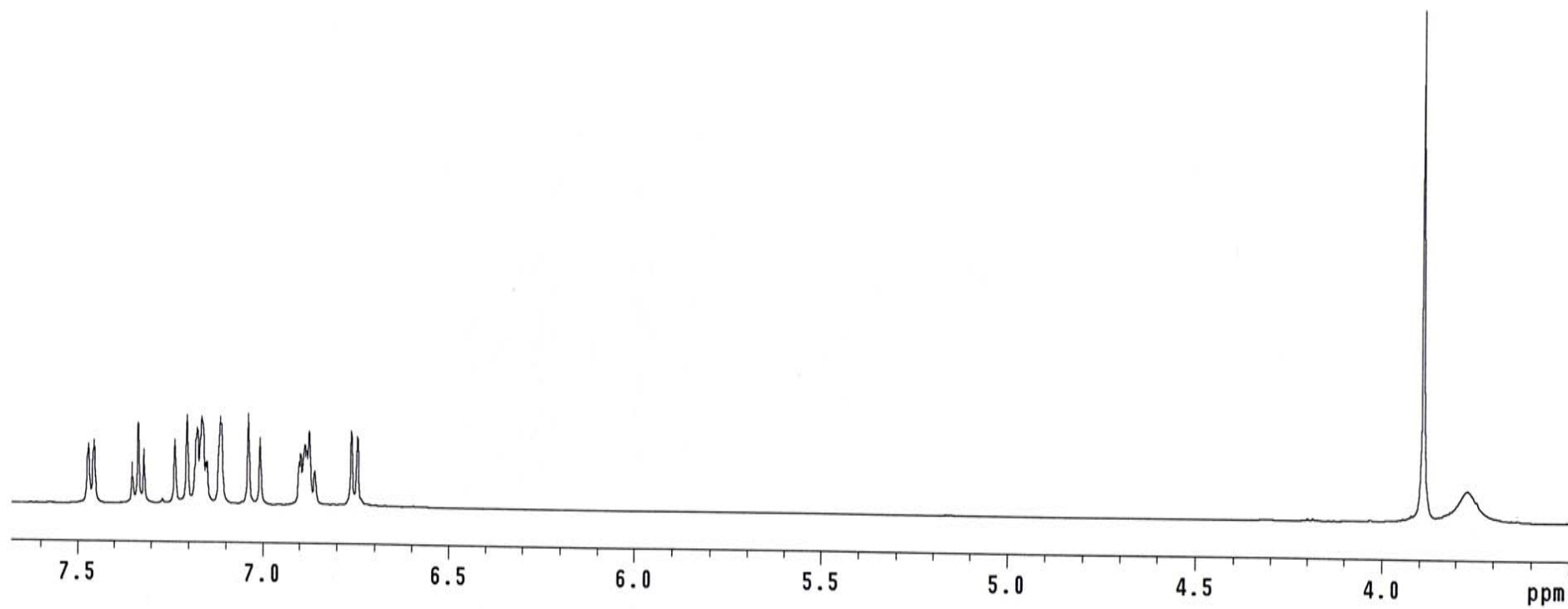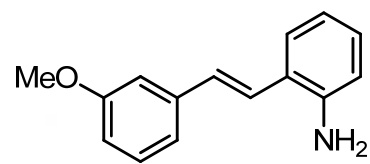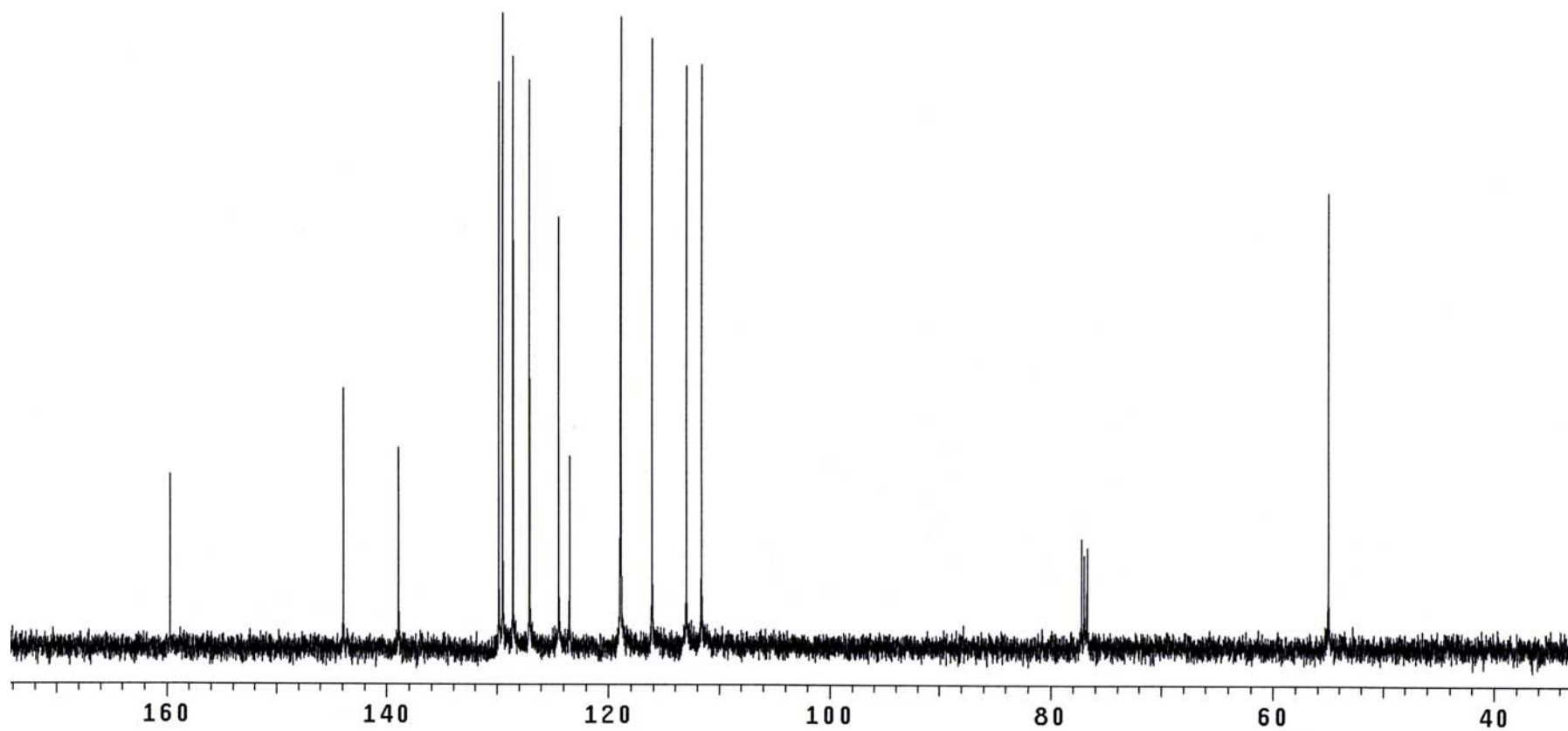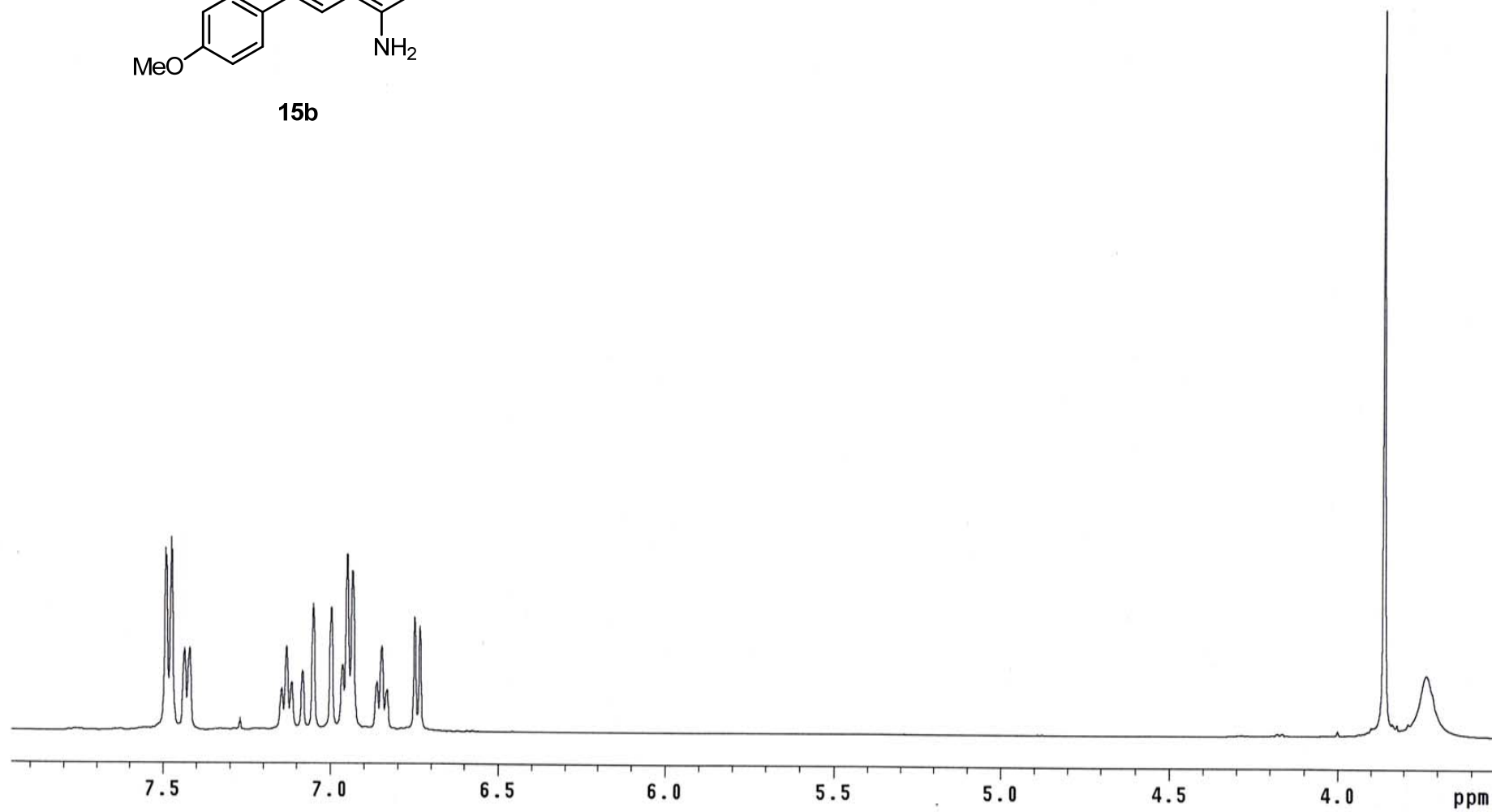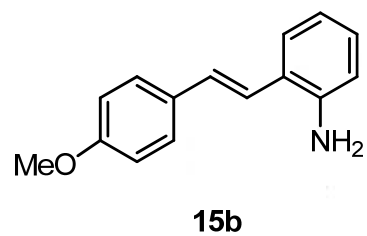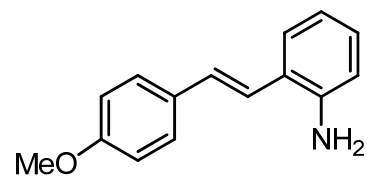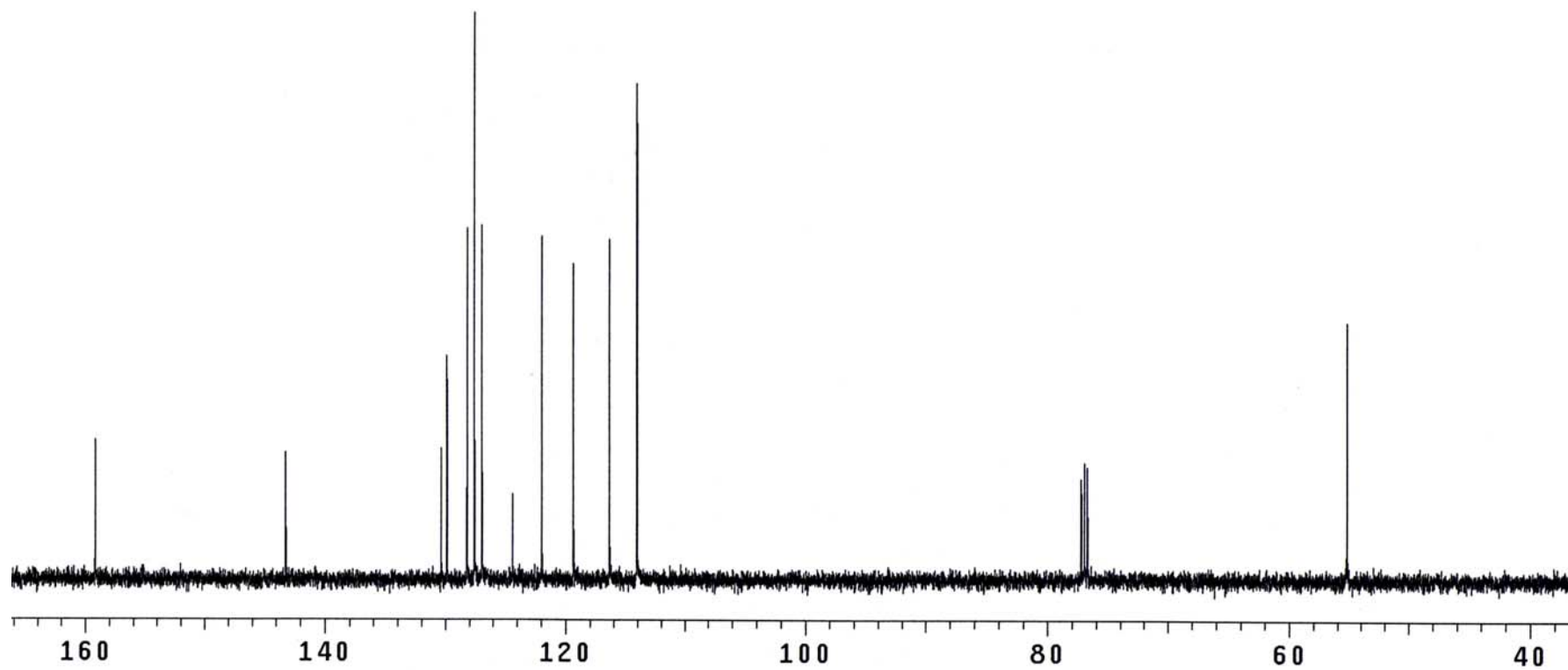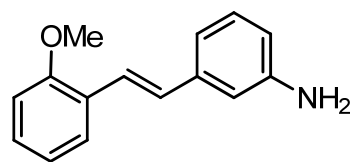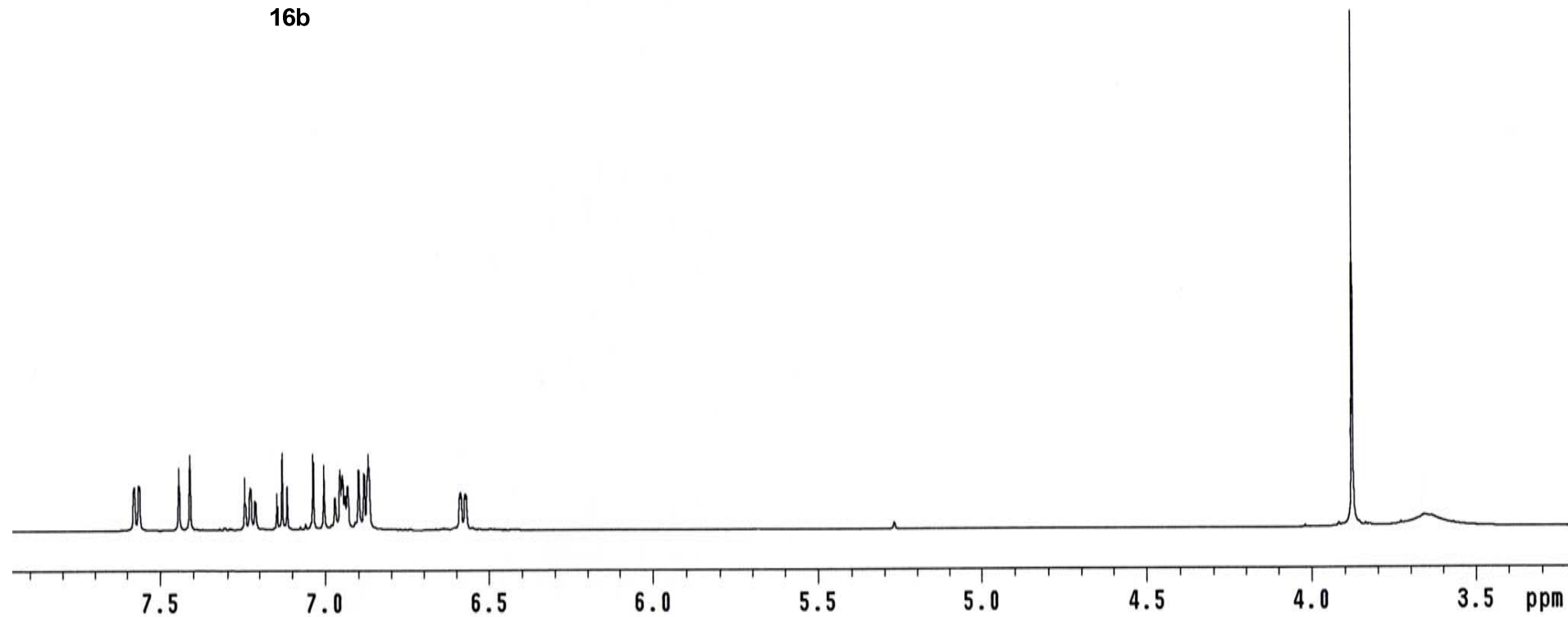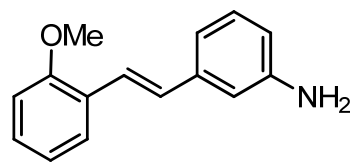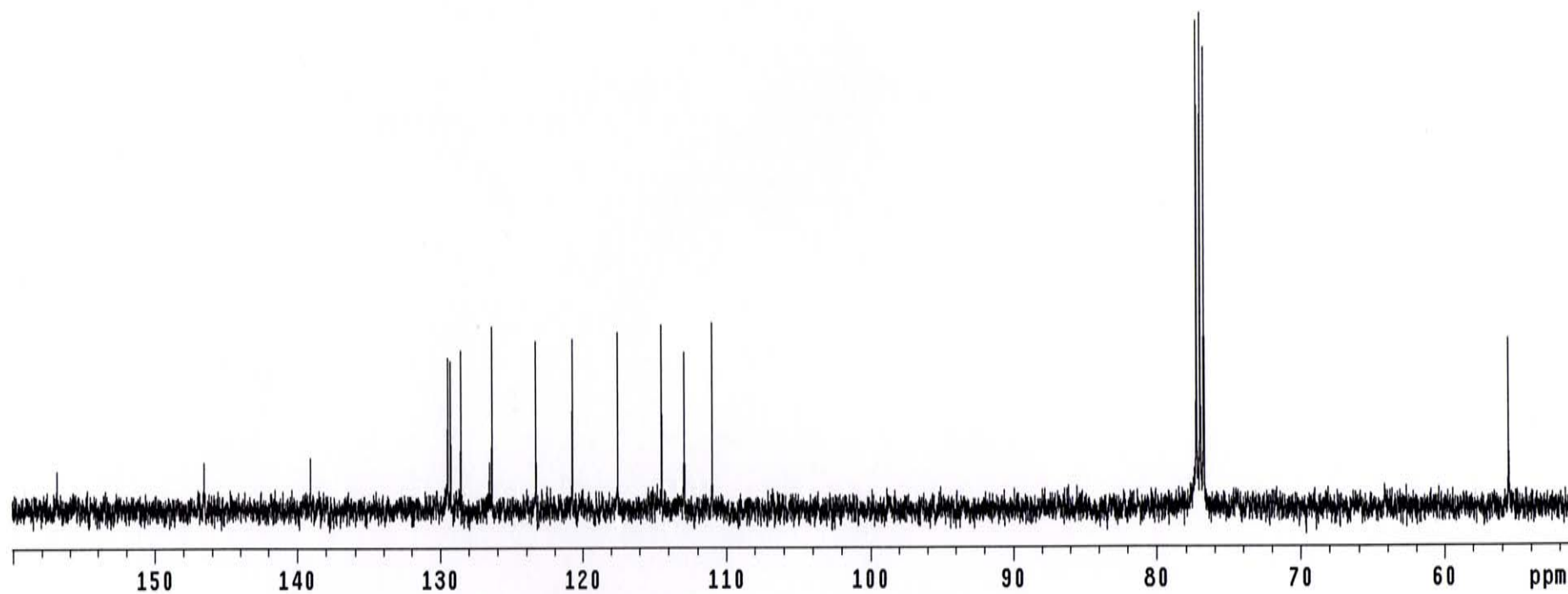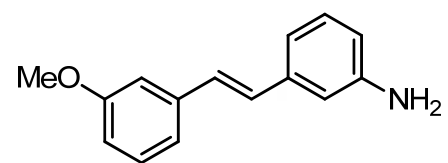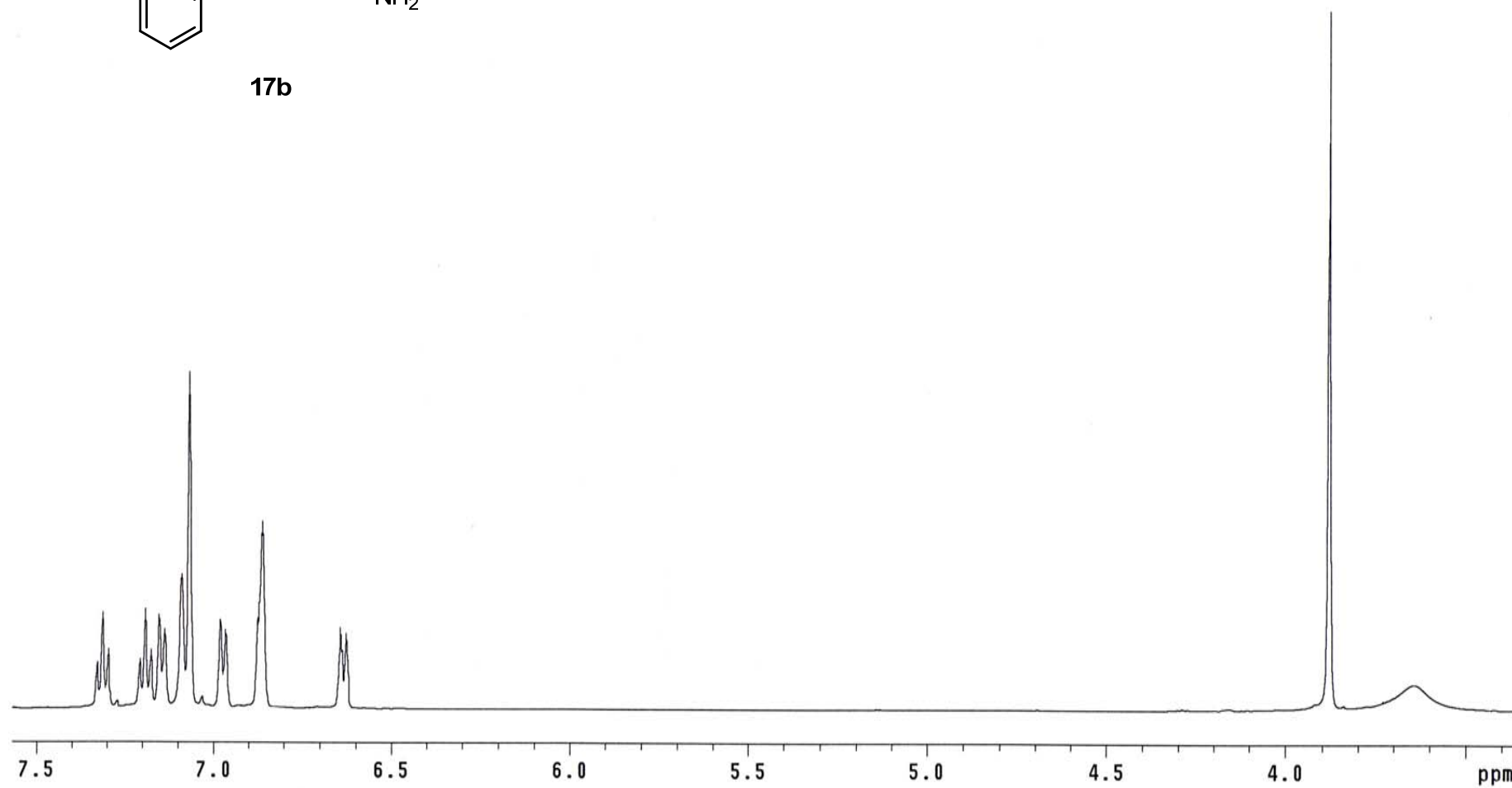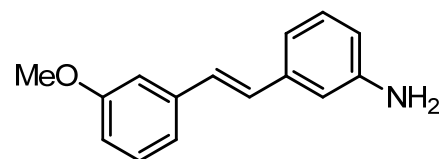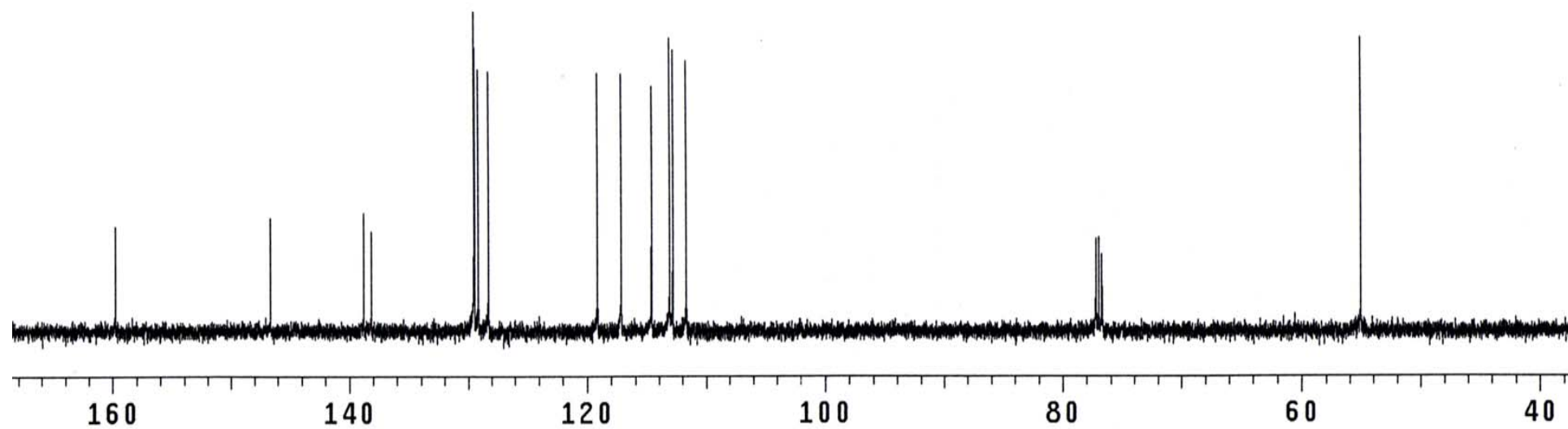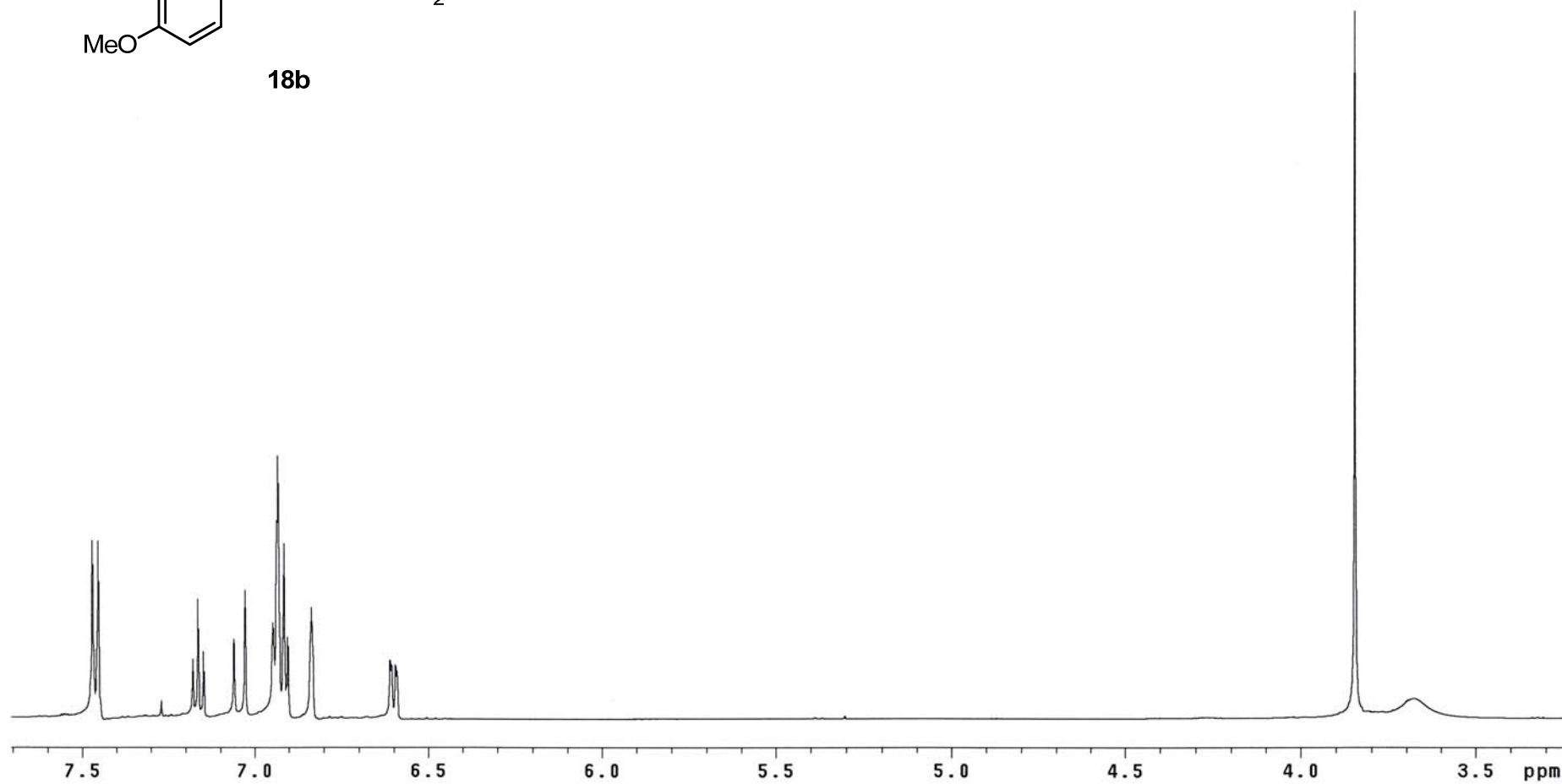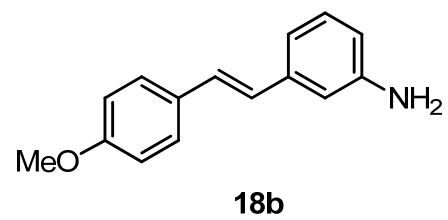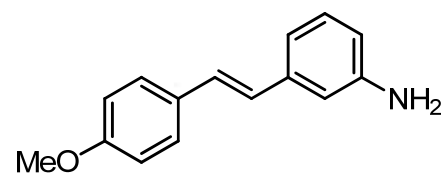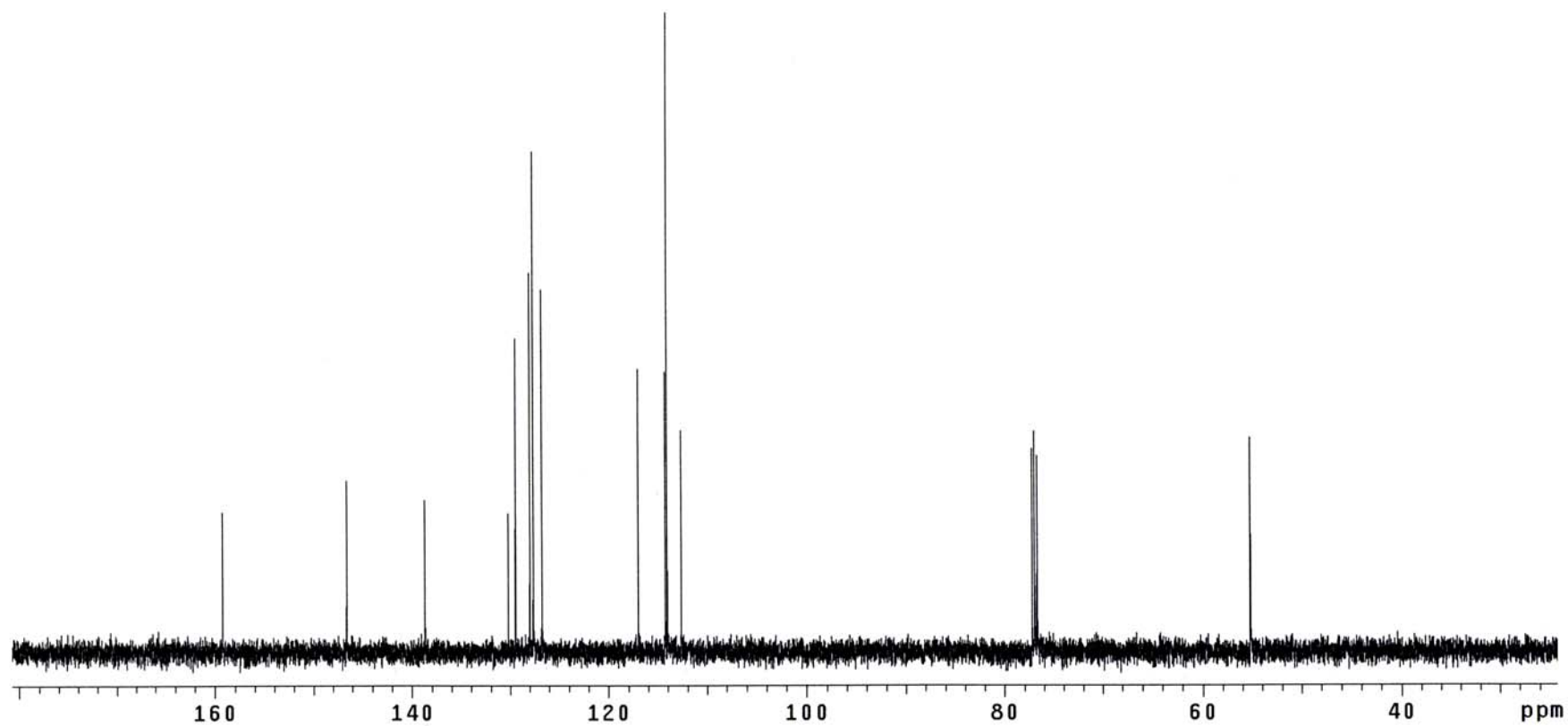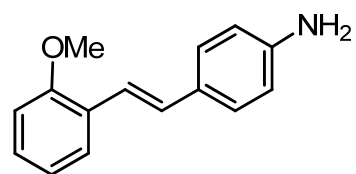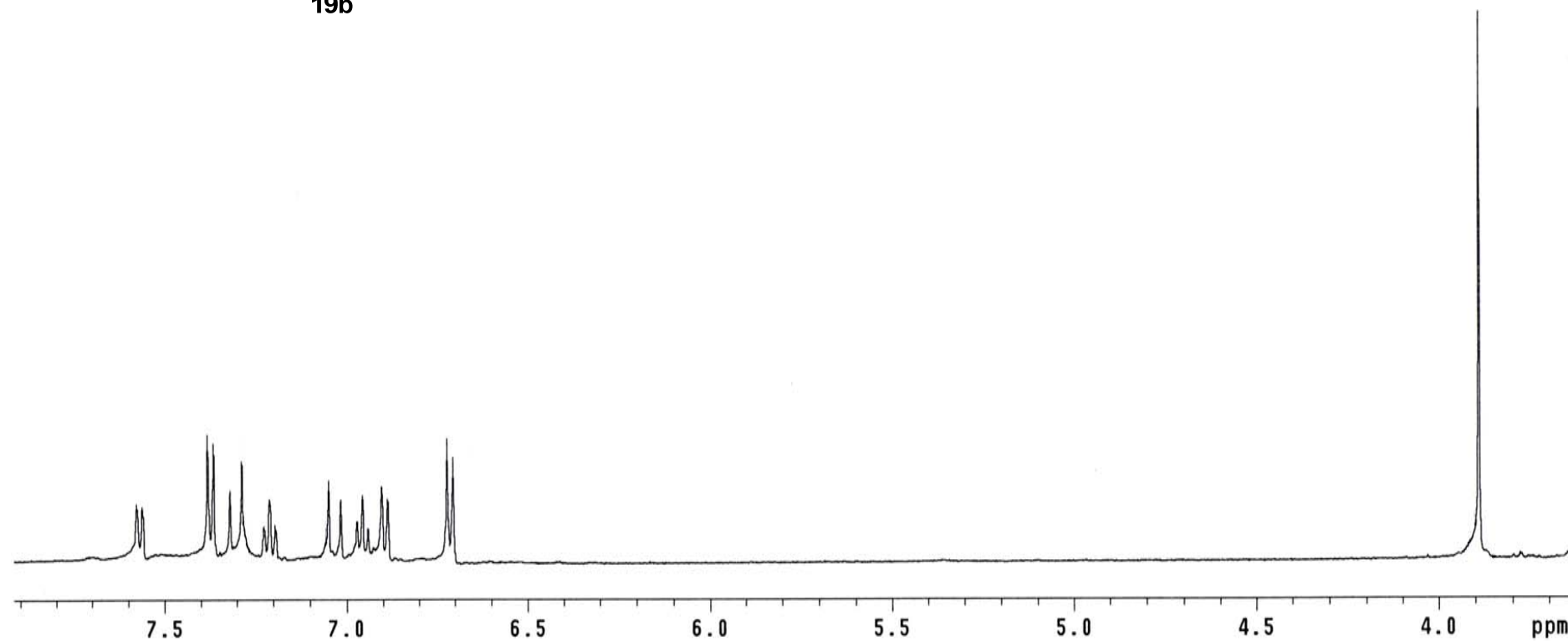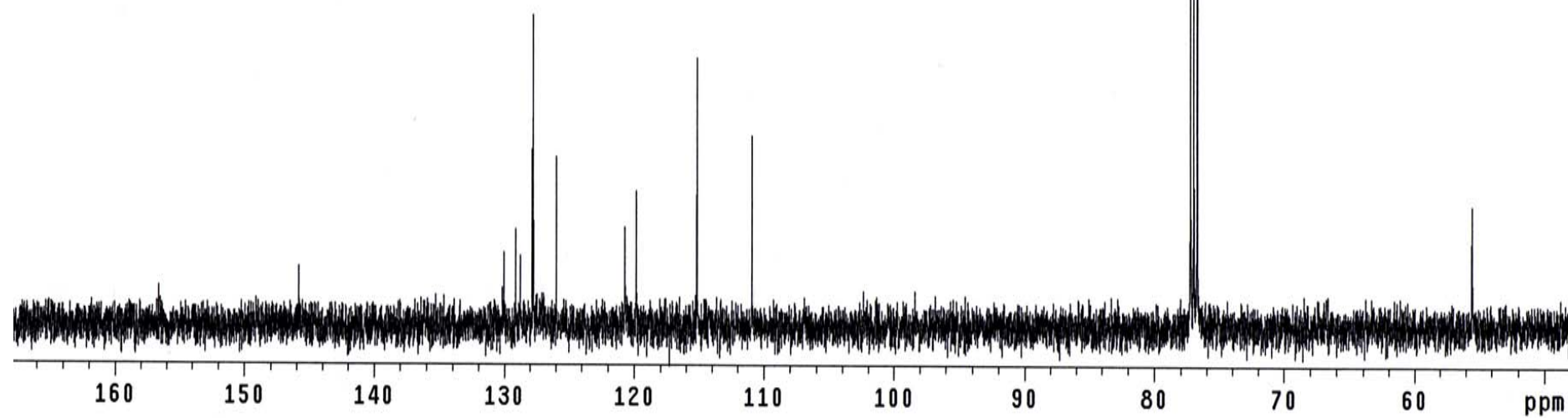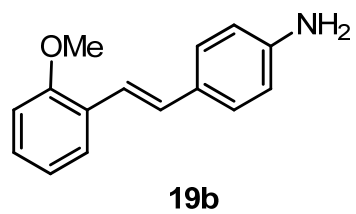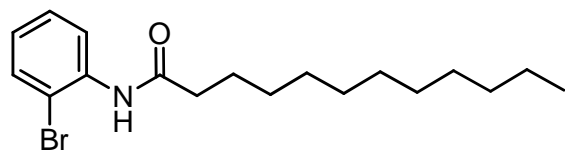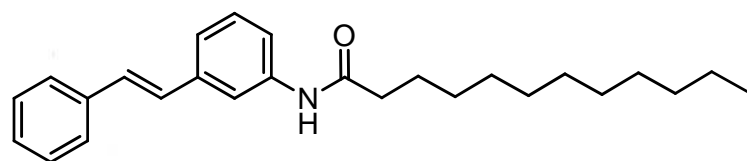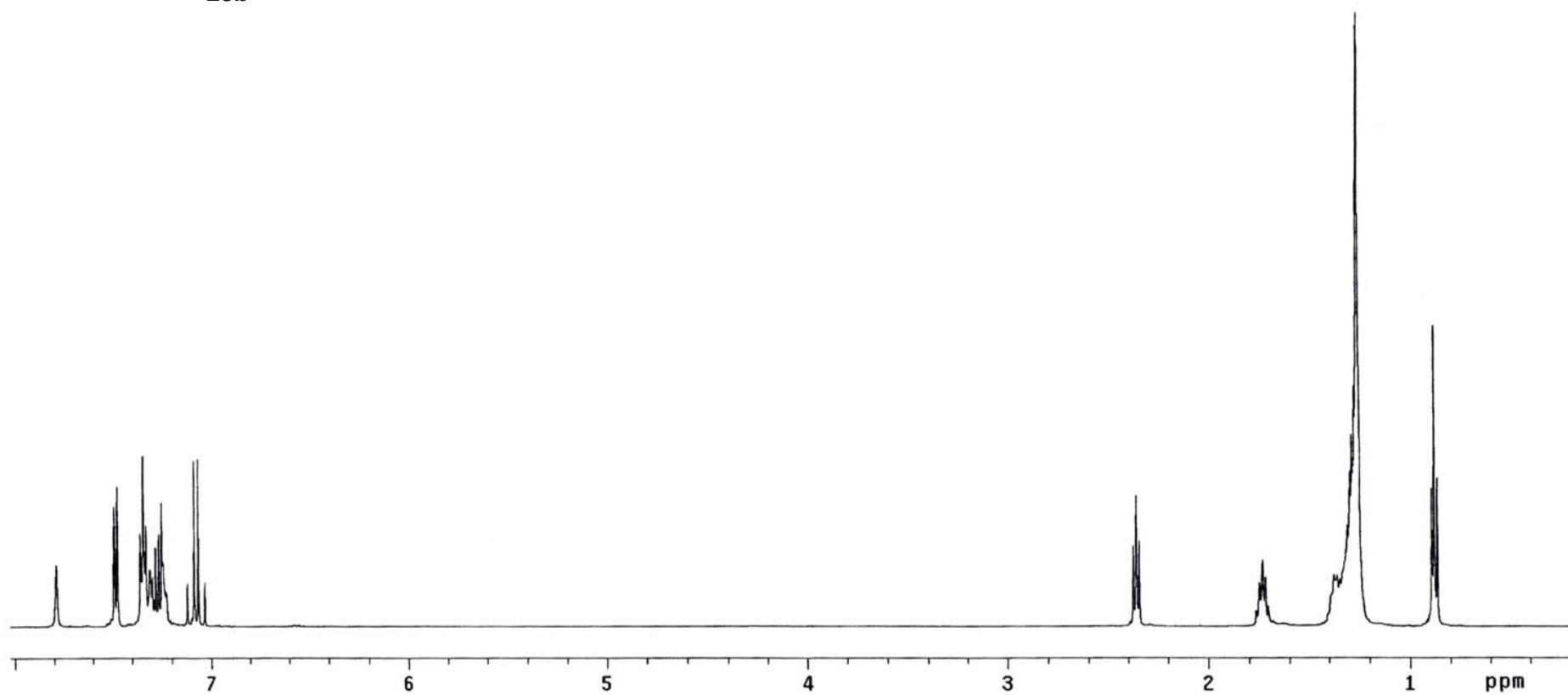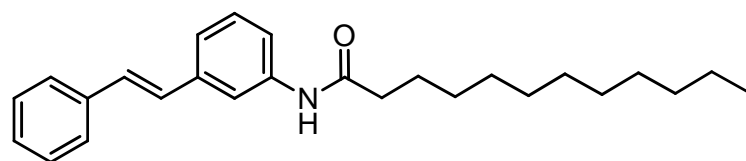
N-(2-bromophenyl)dodecanamide

N-(2-bromophenyl)dodecanamide

22b

22b

23b

23b

24b

24b