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Determination of the Absolute Configuration of Aegelinol by Crystallization of Its Inclusion Complex with β-Cyclodextrin

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Abstract: The absolute configuration and structure of aegelinol, a pyranocoumarin isolated from *Ferulago asparagifolia* Boiss (Apiaceae), has been determined by crystallography. Crystal structure of the inclusion complex of aegelinol in β-cyclodextrin was determined (a = 15.404(1) Å, b = 15.281(1) Å, c = 17.890(1) Å, α = 99.662(1), β = 113.4230(1), γ = 102.481(1)°, P1; R_1 = 6.71%) and allowed unambiguous determination of the absolute configuration of the stereogenic center of aegelinol. The pyranocoumarin guest is included within the cylindrical cavity formed by dimeric β-cyclodextrin molecules with a head-to-head arrangement. Crystal structure of aegelinol alone was also determined (a = 6.8921(3) Å, b = 11.4302(9) Å, c = 44.964(3) Å, P2$_1$2$_1$2$_1$; R_1 = 4.44%) and allowed precise determination of its geometry. Aegelinol crystallizes with three molecules in the asymmetric unit held together by H-bonds and π-stacking interactions.

1. Introduction

Aegelinol (**1**) is a pyranocoumarin that was isolated from *Ferulago asparagifolia* Boiss (Apiaceae), a species growing in Turkey [1]. Aegelinol has also been reported as a minor lactonic constituent of *Aegle marmelos* [2]. Three abundant pyranocoumarins (Figure 1) (grandivittin (**2**), agasyllin (**3**) and aegelinol benzoate (**4**)) were also isolated from the roots of *Ferulago campestris* collected in Sicily and both aegelinol (**1**), their hydrolysis product, and agasyllin (**3**) showed a significant antibacterial effect against Gram-negative and Gram-positive bacteria [3].

Figure 1. The chemical structure and numbering scheme of pyranocoumarins.
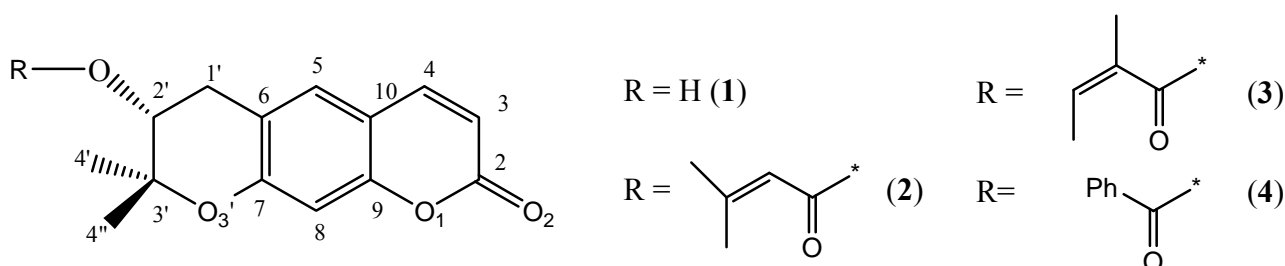
The absolute configuration of **1** could not be determined because NMR spectral data of α and β epimers do not significantly differ. Optical rotation measurements, however, suggest an α configuration for the compound [1]. Compounds **2-4** are ester derivatives of aegelinol (**1**), whose absolute configuration at C-2' was also not clear. Abyshev *et al*. [4] reported an (S) C-2' absolute configuration, although they cited the degradative determination of the absolute configuration performed by Lemmich [5]. On the other hand, Erdelmeyer *et al*. [6] indicated an (R) C-2' configuration. In both cases, the reported optical rotation of the compound was negative. Hydrolysis of the three esters gives the same compound (**1**) with a negative optical rotation. Consequently, Basile *et al*. [3] decided to determinate the absolute stereochemistry by means of Mosher's and Horeau's (indirect) methods and assigned from their NMR data an (R) C-2' absolute configuration for **1** and its esters **2-4**.

Rigorous differentiation of enantiomers by the X-ray diffraction method is now possible through the use of the Flack parameter [7,8]. However, compounds composed only of light atoms (e.g., C, H, O) give rise to standard uncertainties on the Flack parameter which are too large for reliable absolute configuration determination [9].

In many cases, it may thus be necessary to perform a derivatization of the compound to get a crystalline product. If a chiral reagent is used for this purpose, a second stereogenic center of known absolute configuration is introduced and the X-ray structure obtained is then unambiguous and gives

the absolute configuration of the first compound. This approach is classical when salts can be formed, which is not the case for molecules such as aegelinol.

To circumvent this limit, we decided to co-crystallize aegelinol (**1**) with β-cyclodextrin. In principle, formation of such an inclusion complex with a host of known chirality (β-cyclodextrin consists of seven optically active D-glucose units) should allow direct determination of the absolute configuration of the guest (aegelinol). Indeed, in principle, cyclodextrins form diastereoisomers by including optically active guests within their cavity and should be excellent reagents for the resolution of racemic compounds (and subsequent determination of their absolute configuration). In practice, possibly due to their round and symmetrical structure, only low optical resolution is achieved [10]. As a matter of fact, only a limited number of crystal structures of inclusion complexes of cyclodextrins with chiral guests are available in the Cambridge Structure DataBank (CSD), among which permethylated α-cyclodextrin binding S- and R-mandelic acid (CECMAY10 and CECMEC10, respectively) [11] or R- and S-isomers of flurbiprofen and S-ibuprofen (COYXAP10, COYXET20 [12] and RONWOG [13] respectively) included within the cavity of permethylated β-cyclodextrin. Selected examples of crystal structures of cyclodextrin's inclusion complexes with different chiral molecules are presented in Table S1 (supplementary material) [14–32].

In the present work, we report on our efforts to determine by X-ray crystallography the absolute configuration of aegelinol and show that crystallization of an inclusion complex of aegelinol in β-cyclodextrin solved the problem.

2. Results and Discussion

Crystals of aegelinol (**1**) were obtained at room temperature by slow evaporation from a solution in ethanol. The structure belongs to the orthorhombic $P2_12_12_1$ space group with cell parameters $a = 6.8921(3)$ Å, $b = 11.4302(9)$ Å, $c = 44.964(3)$ Å. The structure was solved by direct methods and refined to a final $R_1 = 4.44\%$. Main statistics on data quality and refinement are given in Table 1.

Table 1. Main statistics on data quality and refinement for the crystal structures of aegelinol (**1**) and its inclusion complex in β-cyclodextrin (**1-BCD**).

Crystallographic data	Aegelinol	B-CD-aegelinol
Chemical formula	$C_{14}H_{14}O_4$	$2C_{42}H_{70}O_{35}. 2C_{14}H_{14}O_4.23.5H_2O$
Temperature (K)	150.05(10)	150(2)
Wavelength (Å)	Mo Kα (0.71073)	Cu Kα (1.54179)
Crystal system	Orthorhombic	Triclinic
Space group	$P2_12_12_1$	P1
a (Å)	6.8921(3)	15.404(1)
b (Å)	11.4302(9)	15.281(1)
c (Å)	44.964(3)	17.890(1)
α (°)	90.00	99.662(1)
β (°)	90.00	113.423(1)
γ (°)	90.00	102.481(1)

Table 1. *Cont*.

Crystallographic data	Aegelinol	β-CD-aegelinol
Volume (Å^3)	3542.2(4)	3618.4(5)
Z	12	1
Calculated density (g.cm^{-3})	1.385	1.466
F(0 0 0)	1560	1704
Absorption coefficient (mm^{-1})	0.102	1.148
Crystal description	Long Rod	Plate
Crystal size (mm)	0.2 × 0.2 × 0.4	0.13 × 0.32 × 0.37
Theta range for data collection (°)	3.25- 25.02	2.81–69.59
Limiting indices	$-8 \leq h \leq 6$, $-11 \leq k \leq 13$, $-46 \leq l \leq 53$	$-18 \leq h \leq 14$, $-18 \leq k \leq 18$, $-21 \leq l \leq 21$
Reflections collected/unique	10327/5883	36967/15938
R (int)	0.0246	0.0355
Data/restraints/parameters	5883/0/491	15938/88/2082
Goodness-of-fit (S) on F^2	1.097	1.060
Refinement method	Full-matrix least-squares on F^2	Full-matrix least-squares on F^2
Final R indices [I > 2sigma(I)]	4.44%	6.71%
Final R indices (all data)	5.22%	7.1%
Largest difference peak and hole (e/Å^3)	0.218 and −0.229	1.249 and −0.419
Flack × parameter	0.4(9)	0.5(2)

The structure reveals an infinite three-dimensional network stabilized mainly by ideal intramolecular hydrogen bonds in addition to very strong π-stacking interactions along the *a* direction. Aegelinol crystallizes with three molecules in the asymmetric unit held together by H-bonds and π-stacking interactions (Figure 2). The three molecules are identical with negligible differences in the three mutual RMSDs: A&B = 0.0894, A&C = 0.0642 and B&C = 0.105 Å. This is further confirmed by similar geometrical features (Table 2 and supplementary data).

The coumarin moiety is planar with maximum deviations from the least-squares planes of 0.0247, 0.0396 and 0.0181 Å for the three molecules A, B and C, respectively. The dihedral angles between these planes in crystal packing are AˆA=2.65(3)°, AˆB = 13.39(3)°, BˆC = 2.99(4)° and AˆC =10.76(2)°. The bond lengths and bond angles observed in this structure are comparable with those reported for other coumarin derivatives in CSD (e.g., GEHWEW [33] and MOFJIA [34]).

Figure 2. (**a**) Morphology of crystals of aegelinol (**1**) obtained by slow evaporation from a solution in ethanol; (**b**) Conformation of the three independent molecules in the asymmetric unit showing thermal motion (ORTEP, 30% probability).

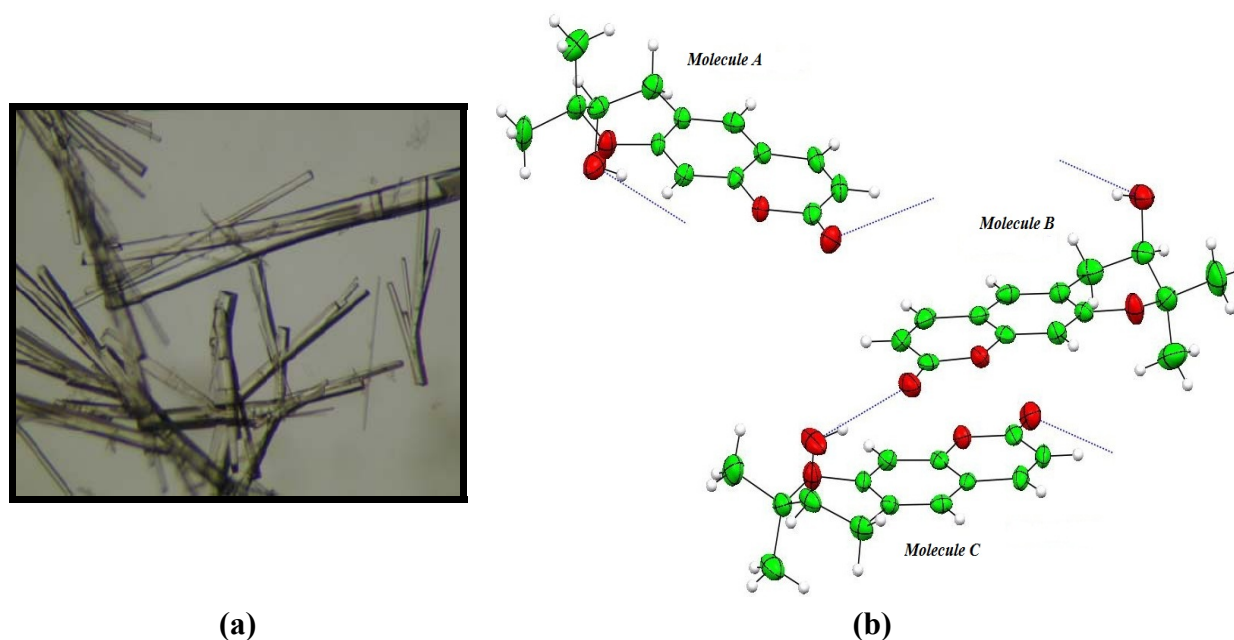(**a**)   (**b**)

Table 2. Selected structural features (bond lengths (Å), valence and torsion angles (°)) of the crystal structures of aegelinol (**1**, three molecules in asymmetric unit (**A, B, C**)) and its inclusion complex in β-cyclodextrin (**1-BCD** two molecules in asymmetric unit (**A, B**))

Geometrical aspects	1 A	1 B	1 C	1-BCD A	1-BCD B
C6-C1`	1.503(3)	1.501(3)	1.501(3)	1.49(2)	1.52(2)
C1`-C2`	1.518(4)	1.523(4)	1.520(4)	1.45(2)	1.49(3)
C2`-O2`	1.431(3)	1.422(3)	1.429(3)	1.43(2)	1.43(2)
C2`- C3`	1.522(4)	1.527(4)	1.521(4)	1.52(2)	1.45(3)
C3`-O3`	1.463(3)	1.470(3)	1.467(3)	1.46(2)	1.50(2)
C7-O3`	1.363(3)	1.361(3)	1.34(2)	1.35(1)	1.36(2)
C7-C6-C1`	120.0(2)	120.0(2)	120.1(2)	121(1)	120(1)
C6-C1`-C2`	110.4(2)	111.6(2)	110.2(2)	110(1)	113(2)
O2`-C2`-C1`	111.4(2)	111.8(2)	111.3(2)	110(1)	114(2)
C1`-C2`-C3`	110.3(2)	110.7(2)	110.6(2)	114(1)	112(2)
O3`-C3`-C2`	109.7 (2)	109.7(2)	110.6(2)	106(1)	114(1)
O3`-C7-C6	122.9(2)	123.0(2)	123.2(2)	122(1)	125(1)
C7-O3`-C3`	118.6 (2)	118.4 (2)	119.0(2)	118(1)	114(1)
C5-C6-C1`-C2`	−156.9(2)	−159.3(2)	−154.0(2)	−165(1)	170(2)
C7-C6-C1`-C2`	21.4(3)	20.5(3)	24.7(3)	14(1)	−10(3)
C6-C1`-C2`-O2`	70.3(3)	79.0(3)	71.9(3)	−172(1)	−78(3)
C6-C1`-C2`-C3`	−50.4(3)	−47.1(3)	−50.8(3)	−46(1)	37(3)
O2`-C2`-C3`-O3`	−62.3(3)	−67.7(3)	−65.7(3)	−173(1)	68(2)
C1`-C2`-C3`-O3`	60.0(3)	58.2(3)	57.7(3)	62(1)	−55(2)

Table 2. *Cont.*

Geometrical aspects	1 A	1 B	1 C	1-BCD A	1-BCD B
O2`-C2`-C3`-C4`	177.6(2)	172.2(2)	173.7(2)	−60(1), 69(2)	−176(1), −53(2)
C1`-C2`-C3`-C4`	−60.0(3)	−62.0(3)	−62.9(3)	175(1), −56(2)	62(2), −175(2)
C1`-C6-C7-O3`	−0.4(4)	−4.0(4)	−4.3(4)	1(1)	0(2)
C8-C7-O3`-C3`	−172.5(2)	−166.6(2)	−172.4(2)	−162.8(9)	162(1)
C6-C7- O3`-C3`	10.1(3)	15.8(3)	11.0(4)	18(2)	−16(2)
C2`-C3`-O3`-C7	−39.6(3)	−42.6(3)	−37.6(3)	−47(1)	44(2)
C4`-C3`-O3`-C7	82.3(3)	79.1(3)	84.9(3)	−165(1), 75(1)	−77(2), 172(2)

The packing of molecules in the unit cell gives some indication of the types of interaction that such a molecule would show in biological systems. Two types of interaction are found. First, the molecules are partially flat and pack in planes, where molecules A and its symmetrical equivalences are stacked one above the other in the opposite direction. B and C molecules have exactly the same arrangement as A molecules. The distances between these planes and the centroid of aromatic ring in the next layer are A-A = 3.481Å and B-C = 3.391Å. This arrangement allows optimal π-π interaction (Figure 3 (see also supplementary material Figure S2)). Second, the carbonyl group interacts with hydroxyl groups in the next layer's molecule with OH···O distances ranging from 2.816(2) to 2.903(3) Å, as shown in Figure 2, Table 3.

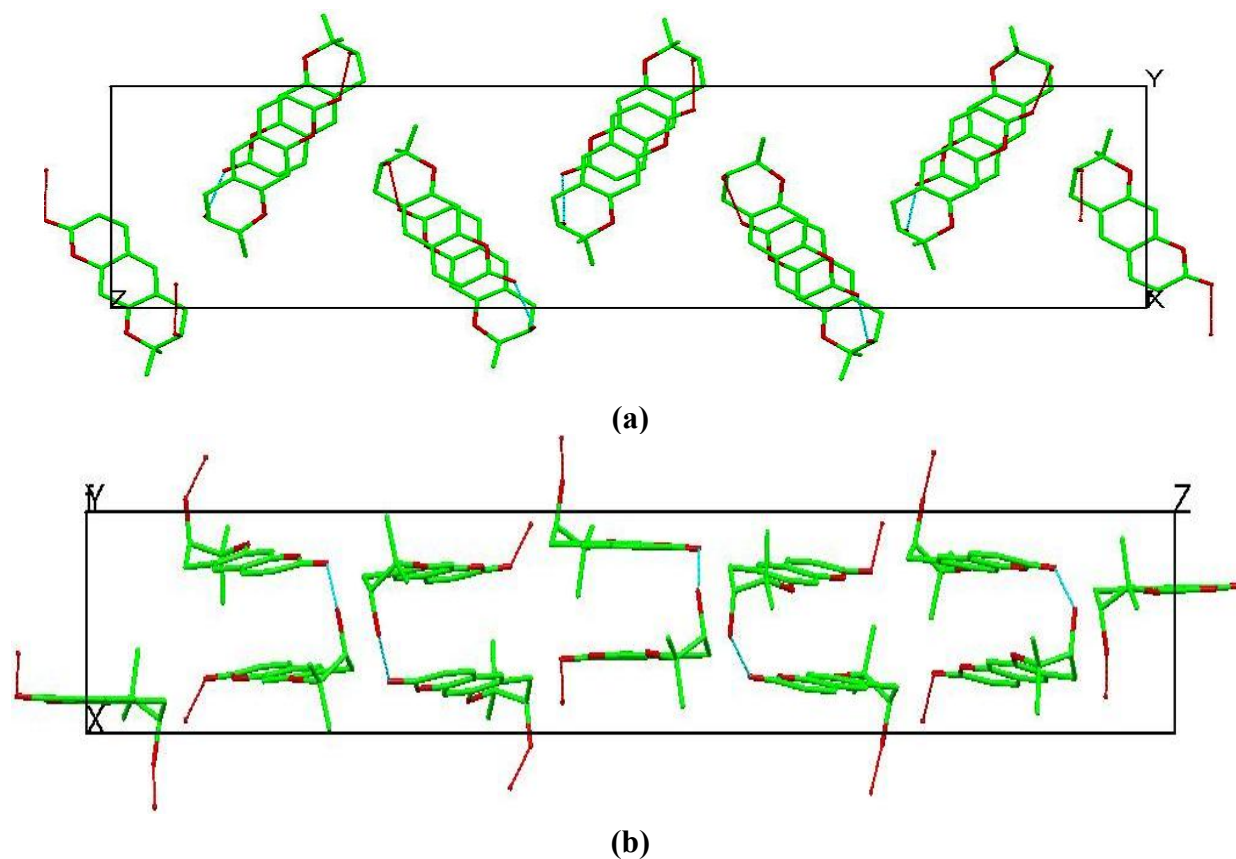**Figure 3.** Crystal packing of aegelinol. (**a**) Stacking along the *a* axis; (**b**) Stacking along the *b* axis.

(a)

(b)

Table 3. Selected hydrogen bonds.

D	A	d (D-H) (Å)	d (H...A) (Å)	d (D...A) (Å)	<(DHA) (°)
Aegelinol (**1**)					
O(2`A)	O(2A) *i*	0.84	2.08	2.902(3)	164.5
O(2`B)	O(2C) *ii*	0.84	2.01	2.816(2)	159.4
O(2`C)	O(2B)	0.84	2.03	2.851(3)	165.8
BCD-1 complex (**1-BCD**)					
O2`A1	O(13W)	0.84	2.122	2.73(1)	129.16
O2`B1	O(61A)	0.84	2.524	3.278(3)	150.01
O2`A2	O(17`W)	0.84	2.04	2.49(2)	145.6
O2`B2	O(63B) *iii*	0.84	2.065	2.83 (2)	151.37
O(65B)	O2`A2 *iv*	0.84	2.05	2.76(2)	144.5

Symmetry transformations used to generate equivalent atoms: *i* *[x-1/2, -y+3/2, -z+2]*; *ii* *[x+1, y, z]*;

iii *[x+1, y, z+1]*; *iv* *[x, y, z+1]* (See supplementary data for molecular numbering Figure S1 and Figure S3).

Statistics of the Flack parameters (x = 0.4(9)) confirm that the absence of suitable heavy atoms (anomalous scatterers) in the molecule is a limitation for reliable absolute configuration of such molecules by X-ray diffraction [9].

Crystal structure of the inclusion complex of aegelinol in β-cyclodextrin (**1-BCD**) was also determined (*a* = 15.404(1) Å, *b* = 15.281(1) Å, *c* = 17.890(1) Å, *α* = 99.662(1)°, *β* = 113.423(1)°, *γ* = 102.481(1)°, P1; R_1 = 6.71%) and allowed unambiguous determination of the absolute configuration of the stereogenic center of aegelinol. Crystals were grown from a solution in water/ethanol (50:50). Two morphologies of crystals are observed: the long and yellow crystals correspond to aegelinol alone, and parallelepiped and colorless ones are related to the inclusion complex with β-cyclodextrin. Primary statistics on data quality and refinement are given in Table 1. Figure 4 presents the structure of the inclusion complex. From this structure, the absolute configuration of C2' of aegelinol can unambiguously be assigned and corresponds to the (R) enantiomer.

Interestingly, disorder of the included aegelinol molecule within the cyclodextrin cavity was modeled in two sites, nearly similar, with 60:40 occupancy. The coumarin moieties are planar with maximum deviations from the least-squares planes of 0.0034 and 0.0038 Å for the two molecules A and B, respectively. The dihedral angles between these planes, inside the cavity, is 22.2(4)° in the first site and 3.0(3)° in the second site.

The solution of the crystal structure revealed a 2:2 stoichiometry. The pyranocoumarin guest is included within the cylindrical cavity formed by dimeric β-cyclodextrin molecules with a head-to-head arrangement. Crystalline complexes of hydrogen-bonded β-cyclodextrin dimers complexed with organic guest molecules have been found previously to adopt one of four crystal packing motifs: a channel motif (***Ch***), a checker board motif (***CB***), an intermediate motif (***Im***), and, most recently, a tetrad motif (***Tt***). Packing diagrams are available in different reports on β-cyclodextrin complexes; the reader is referred to these works for both diagrams and discussion of the different structural properties for these packing arrangements [35–37]. The CD host molecules are associated through hydrogen bonding between secondary hydroxyl groups as dimers, packing in the previously observed (***Im***)

packing arrangement [35–37]. The fact that this arrangement is common to these complexes of β-CD is important for two reasons:

(1) It implies that crystal packing does not severely limit conformational and translational freedom of guest molecules.

(2) It provides an environment for the guest molecules that resembles a macromolecular binding pocket by providing a hydrophobic pocket rimmed with a scaffolding of hydrophilic binding sites that can interact directly with guest molecules or via bridging water molecules.

Figure 4. (**a**) Morphology of crystals of aegelinol-β-cyclodextrin inclusion complex (**1-BCD**). Two forms of crystals are observed: (**I**) corresponds to cell parameters of aegelinol alone, and (**II**) corresponds to cell parameters of the inclusion complex; (**b**) Two perpendicular views showing the conformation of the **1-BCD** inclusion complex (ORTEP, 30% probability) and confirming the R configuration at C2'. Only one component of the disorder of the guest molecule inside the cyclodextrin cavity is presented and hydrogen atoms and water molecules are removed for clarity; (**c**) View of the two aegelinol molecules after removing all the rest of the structure to show their configuration.
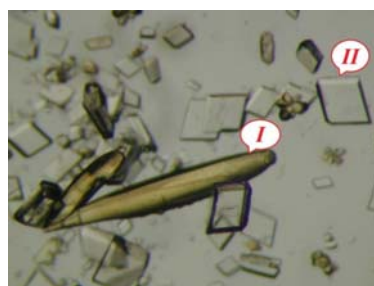

(a)
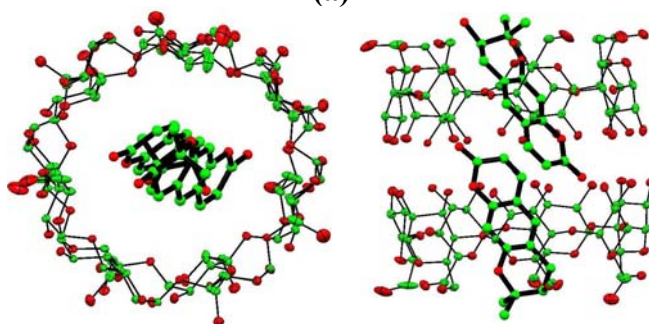
(b)
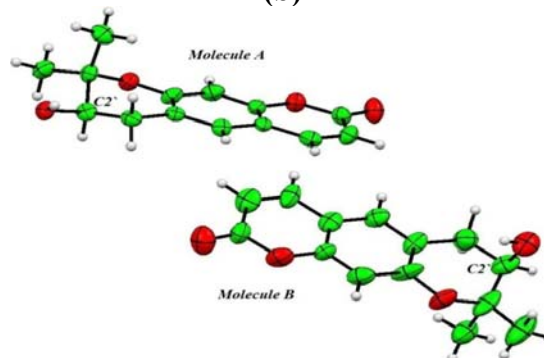
(c)

As a consequence, guest molecules experience considerable variation in their specific interactions with the host molecules and waters of hydration, often resulting in the observation of crystallographic disorder. This semichannel-type packing orients along the c-axis direction (Figure 5).

Figure 5. Crystal packing of β-cyclodextrins in the **1-BCD** crystal structure shown along the *a*, *b* and *c* directions. Guest molecules (aegelinol) and waters are omitted for clarity.
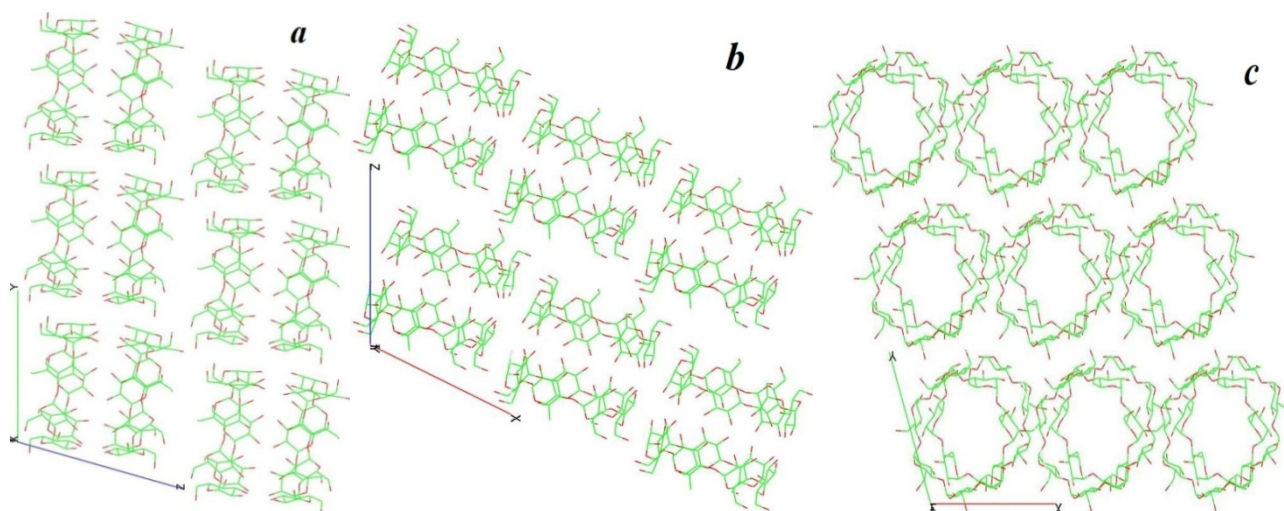
One hydrogen bond is observed between the cyclodextrin and the first site and two in the second site of the disordered guest Table 3 (See supplementary material Figures S1 and S3). These hydrogen bonds are between the hydroxyl groups on the guest and on the primary side of cyclodextrin. The two molecules of aegelinol stack together (mainly involving the coumarin moieties) and are stabilized by hydrophobic interactions within the cavity formed by two cyclodextrins (Figure 4). The distances between the planar coumarin and the centroid of the other coumarin are 3.233Å in the first site and 3.215 Å in the second site. Moreover, this arrangement serves a strong π-π interaction (See supplementary material Figure S2). Notable differences are observed at the level of the pyroano ring of the two aegelinol molecules (RMSD = 0.871 Å and Table 2) most likely resulting from less constraints in **1**-BCD compared to the structure of **1** alone. Twenty-four ordered water molecules are also observed in the structure. They are situated outside the cavity at the borders of the toroid rims and in interstices between β-CD molecules. Five water molecules are disordered over two sites.

3. Experimental Section

Isolation and purification of aegelinol (**1**) has been reported elsewhere [1]. Briefly, roots of *F. asparagifolia* (500 g) were dried, powdered and extracted successively with light petroleum, EtOAc and MeOH at room temperature. Extracts were then evaporated and lyophilized. The EtOAc extract (4.0 g) was chromatographed over silica gel columns. Six compounds were isolated from the EtOAc extract, one of which corresponding to aegelinol.

Crystals of aegelinol were prepared by dissolving a 1–2 mg amount of aegelinol in a minimal amount of ethanol. The mixture was then stored at room temperature. Colorless long crystals suitable for X-ray data collection were obtained by slow evaporation after one week.

Crystals of the β-CD in complex with aegelinol were prepared by dissolving a few milligrams of β-CD in water (2 mL) at 65 °C and aegelinol in ethanol (2 mL) at 65 °C, in stoichiometric (2:1) ratio with respect to β-CD. Both solutions were mixed and stirred at 65 °C for 6 hours. Then the mixture was stored at room temperature. Two forms of crystals (colorless plates and pale yellow needle) suitable for X-ray data collection were obtained by slow evaporation after one week.

Single crystals of aegelinol and of the β-CD-aegelinol complex were mounted on the head of a 4-circle kappa goniometer. X-ray diffraction experiment was carried out at low temperature (150 K) on an Oxford Gemini R Ultra using Mo Kα (0.71073 Å) (aegelinol) or Cu Kα (1.54179 Å) radiation (inclusion complex). Data reduction was carried out with the CrysAlisPro program [38]. Crystal data collection and refinement details are listed in Table 1.

The structure of aegelinol was determined by direct methods and refined using full-matrix least-squares based on F^2 with the program SHELX [39], based on 491 parameters and 0 restraints. The refinement converged to $R_1 = 4.44\%$ for $I > 2\sigma(I)$ and 5.22% for all data. Hydrogen atoms were calculated at ideal positions and refined using a riding model.

The structure of the β-CD-aegelinol complex was solved by atomic coordination replacement methods using a β-CD complex structure from CSD [BOSZOZ] [40] with the same cell parameters after making suitable transformation and deleting the guest from the structure. Extra electronic density in the cavity corresponded to aegelinol. This structure was refined using full-matrix least-squares on F^2 with the program SHELX [39], based on 2083 parameters and 89 restraints. The refinement converged to $R_1 = 6.71\%$ for $I > 2\sigma(I)$ data and 7.10% for all data. Most hydrogen atoms were calculated at ideal positions and refined using a riding model and the rest of the hydrogen atoms were located on a different Fourier map. Ordered water molecules were located in a difference Fourier map. All non-hydrogen atoms were anisotropically refined. Secondary hydroxyl groups were refined with two alternate conformations. Disorder of the included aegelinol molecules within the cyclodextrin cavity, five disordered water molecules and two primary hydroxyl groups in β-CD were modeled in two sites. The second sites were located from difference Fourier maps.

Coordinates have been deposited at the Cambridge Structure DataBank (CCDC codes 884098 and 884099) and are available as supplementary material to this article.

4. Conclusions

Crystals of aegelinol (**1**) alone were obtained and led to determination of its crystal structure. Statistics of the Flack parameters confirm that the absence of suitable heavy atoms (no strong anomalous scatterers) in the molecule is a limitation for reliable absolute configuration of such molecules by X-ray diffraction. Derivatization of the molecule by formation of a chiral salt was not an alternative as the molecule does not contain any basic or acidic function. As an alternative, we were able to grow crystals of the inclusion complex of aegelinol in β-cyclodextrin and refined the crystal structure of the complex, which allowed unambiguous determination of the absolute configuration of the stereogenic center of aegelinol. The (R) configuration of C2' obtained by this crystallographic study confirms previous conclusions based on indirect NMR data.

Acknowledgments

Conflict of Interest

The authors declare no conflict of interest.

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