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(1α,8β)-6β-Benzoyloxy-6-dehydroxyheteratisine from *Aconitum zeravschanicum*

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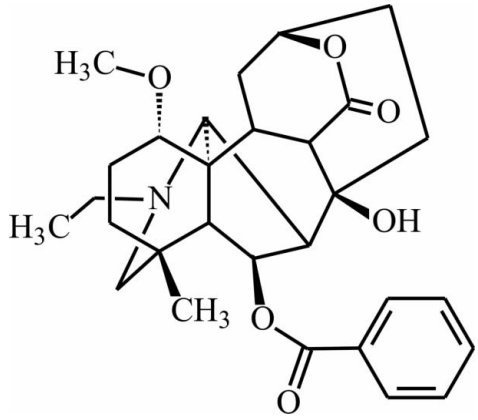
Key indicators: single-crystal X-ray study; T = 300 K; mean σ(C–C) = 0.009 Å; R factor = 0.067; wR factor = 0.142; data-to-parameter ratio = 7.5.

The title compound, $C_{29}H_{37}NO_6$, was isolated from *Aconitum zeravschanicum* and exhibits antiarhythmic activity. It is a derivative of the diterpenoid alkaloid heteratisine and as such the core framework of the molecule contains four six-membered, three seven-membered and one five-membered ring. The chair conformation of one of the methoxy-substituted six-membered rings is different from that observed in heteratisine hydrobromide monohydrate. In the latter case, this ring adopts a boat conformation due to a stabilizing intramolecular N—H···O hydrogen bond. In the crystal structure of the title compound, there is only one acidic H atom. This hydroxyl group forms an intermolecular O—H···O hydrogen bond that links molecules into infinite chains along the b axis.

Related literature

For the isolation and idenfication of 6-benzoylheteratisine, see: Aneja *et al.* (1973), Jacobs *et al.* (1947), Nigmatullaev *et al.* (2000). For antiarhythmic activity, see: Salimov *et al.* (1996). For the structure of heteratisine hydrobromide monohydrate, see: Przybylska (1965).



Experimental

Crystal data

$C_{29}H_{37}NO_6$
M_r = 495.60
Orthorhombic, $P2_12_12_1$
a = 10.039 (5) Å
b = 14.107 (8) Å
c = 17.512 (6) Å
V = 2480 (2) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 0.09 mm^{-1}
T = 300 K
0.50 × 0.30 × 0.15 mm

Data collection

Stoe Stadi-4 four-circle diffractometer
Absorption correction: none
2481 measured reflections
2481 independent reflections
1667 reflections with $I > 2\sigma(I)$
3 standard reflections
every 200 reflections
intensity decay: 6.8%

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.067
$wR(F^2)$ = 0.142
S = 1.22
2481 reflections
330 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.22 e Å^{-3}
$\Delta\rho_{min}$ = −0.21 e Å^{-3}

Table 1
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
O3—H3···O1^{i}	0.82	2.25	3.056 (8)	166

Symmetry code: (i) $-x+1, y-\frac{1}{2}, -z+\frac{1}{2}$.

Data collection: *STADI4* (Stoe & Cie, 1997); cell refinement: *STADI4*; data reduction: *X-RED* (Stoe & Cie, 1997); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXTL* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: ZL2222).

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supplementary materials

Acta Cryst. (2009). E**65**, o1682-o1683 [**doi:10.1107/S1600536809023873**]

(1*α*,8*β*)-6*β*-Benzoyloxy-6-dehydroxyheteratisine from *Aconitum zeravschanicum*

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Comment

The title compound of this study, 6-benzoylheteratisine, $C_{29}H_{37}NO_6$, was first obtained synthetically in 1973 (Aneja *et al.*, 1973) and found to be a derivative of a naturally occuring compound (Jacobs *et al.* 1947). Later it was isolated from *Aconitum zeravschanicum* Steinb (Nigmatullaev *et al.* 2000). 6-Benzoylheteratisine exibits antiarhythmic activity that exceeds other antiarrhythmic drugs of the quinidine groups (Salimov *et al.* 1996). The crystal structure of the parent compound was previously established as a salt in the form of heteratisine hydrobromide monohydrate (Przbylslka, 1965).

The molecular structure of the title compound is shown in Fig. 1. The heteratisine skeleton contains four six-membered rings, (***A***, ***C***, ***D*** and ***F***), one five-membered ring (***B***), and three seven-membered rings (e.g. ***E***, others not labeled for clarity) (Fig. 2). Ring ***B*** has an envelope and ring ***C***a more or less regular chair conformation. Ring ***F*** shows a significant distortion and rings ***D*** and ***E*** adopt a boat conformations. The chair conformation of ring ***A*** in the title molecule is different from that observed in heteratisine hydrobromide monohydrate (Przybylska, 1965). For the salt of the parent compound ring ***A*** adopts a boat conformation due to a stabilizing intramolecular N—H···O hydrogen bond between the protonated amine towards the oxygen atom, an interaction not present in the title compound.

The aromatic ring and the acyl-group are rotated against each other, the dihedral angle of their respective planes is 32.6 (9)°. There is only one acidic hydrogen atom in the crystal structure of the title compound. This hydroxyl group forms an intermolecular O—H···O hydrogen bond that links the molecules into infinite chains along the *b*-axis. (Table 1; Fig.3)

Experimental

The title compound was isolated from the chloroform fraction of the leaves of *Aconitum zeravschanicum* by a known method (Nigmatullaev *et al.*, 2000). Crystals suitable for X-ray analysis were obtained by slow evaporation of an ethanol solution at room temperature (m.p. 485–487 K).

Refinement

The hydroxyl H atom was located in a difference Fourier map but was ultimately placed geometrically (with an O—H distance of 0.82 Å). The H atoms bonded to C atoms were placed geometrically (with C—H distances of 0.98 Å for CH; 0.97 Å for CH_2; 0.96 Å for CH_3; and 0.93 Å for C_{ar}) and included in the refinement in a riding motion approximation with U_{iso}=1.2U_{eq}(C) [U_{iso}=1.5U_{eq}(C,O) for methyl and hydroxyl H atoms].

Figures

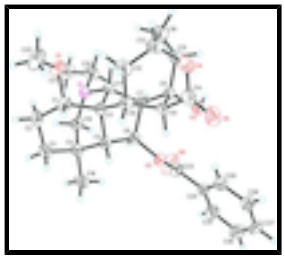

Fig. 1. The molecular structure of 6-benzoylheteratisine, showing the atomic numbering scheme and displacement ellipsoids drawn at the 30% probability level.

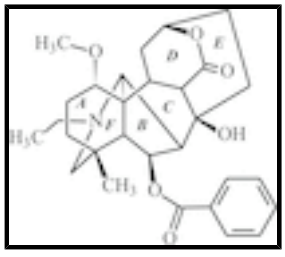

Fig. 2. Ring assigments in 6-benzoylheteratisine.

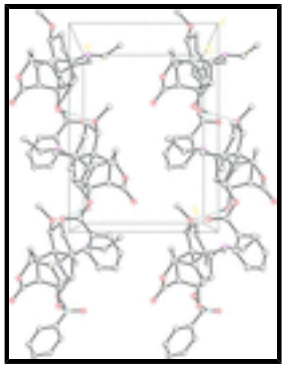

Fig. 3. View along the *c* direction of the crystal packing of 6-benzoylheteratisine, showing the formation of hydrogen bonds (dashed lines). H-atoms not involved in hydrogen bonding have been removed for clarity.

(1α,8β)-6β-Benzoyloxy-6-dehydroxyheteratisine

Crystal data

$C_{29}H_{37}NO_6$	D_x = 1.327 Mg m^{-3}
M_r = 495.60	Melting point: 486(2) K
Orthorhombic, $P2_12_12_1$	Mo $K\alpha$ radiation, λ = 0.71073 Å
Hall symbol: P 2ac 2ab	Cell parameters from 30 reflections
a = 10.039 (5) Å	θ = 10–20º
b = 14.107 (8) Å	μ = 0.09 mm^{-1}
c = 17.512 (6) Å	T = 300 K
V = 2480 (2) Å^3	Prizmatic, colourless
Z = 4	0.50 × 0.30 × 0.15 mm
F_{000} = 1064	

Data collection

Stoe Stadi-4 four-circle diffractometer	R_{int} = 0.0000

Radiation source: fine-focus sealed tube	θ_{max} = 25.0º
Monochromator: graphite	θ_{min} = 1.9º
T = 300 K	h = 0→11
Scan width (ω) = 1.56 – 1.68, scan ratio 2θ:ω = 1.00 I(Net) and sigma(I) calculated according to Blessing (1987)	k = 0→16
Absorption correction: none	l = 0→20
2481 measured reflections	3 standard reflections
2481 independent reflections	every 200 reflections
1667 reflections with $I > 2\sigma(I)$	intensity decay: 6.8%

Refinement

Refinement on F^2	Hydrogen site location: inferred from neighbouring sites
Least-squares matrix: full	H-atom parameters constrained
$R[F^2 > 2\sigma(F^2)]$ = 0.067	$w = 1/[\sigma^2(F_o^2) + (0.0342P)^2 + 1.8083P]$ where $P = (F_o^2 + 2F_c^2)/3$
$wR(F^2)$ = 0.142	$(\Delta/\sigma)_{max}$ = 0.001
S = 1.22	$\Delta\rho_{max}$ = 0.22 e Å^{-3}
2481 reflections	$\Delta\rho_{min}$ = −0.20 e Å^{-3}
330 parameters	Extinction correction: SHELXL97 (Sheldrick, 2008), $Fc^*=kFc[1+0.001xFc^2\lambda^3/\sin(2\theta)]^{-1/4}$
Primary atom site location: structure-invariant direct methods	Extinction coefficient: 0.0056 (10)
Secondary atom site location: difference Fourier map	

Special details

Geometry. All e.s.d.'s (except the e.s.d. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell e.s.d.'s are taken into account individually in the estimation of e.s.d.'s in distances, angles and torsion angles; correlations between e.s.d.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell e.s.d.'s is used for estimating e.s.d.'s involving l.s. planes.

Refinement. Refinement of F^2 against ALL reflections. The weighted R-factor wR and goodness of fit S are based on F^2, conventional R-factors R are based on F, with F set to zero for negative F^2. The threshold expression of $F^2 > \sigma(F^2)$ is used only for calculating R-factors(gt) *etc*. and is not relevant to the choice of reflections for refinement. R-factors based on F^2 are statistically about twice as large as those based on F, and R- factors based on ALL data will be even larger.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (Å^2)

	x	y	z	U_{iso}*/U_{eq}
O1	0.6593 (4)	0.5430 (3)	0.3119 (3)	0.0562 (12)
O2	0.6307 (4)	0.1410 (3)	0.4078 (2)	0.0416 (10)
O3	0.6046 (4)	0.1087 (3)	0.2572 (2)	0.0520 (12)
H3	0.5411	0.0905	0.2317	0.062*
O4	0.8674 (5)	0.2617 (4)	0.1608 (3)	0.0634 (13)
O5	0.9303 (5)	0.1410 (4)	0.2317 (3)	0.0797 (17)

O6	0.4463 (5)	0.0591 (4)	0.4323 (4)	0.099 (2)
N1	0.4096 (5)	0.4035 (4)	0.3632 (3)	0.0437 (13)
C1	0.6830 (6)	0.4820 (4)	0.3766 (4)	0.0473 (17)
H1A	0.7785	0.4850	0.3879	0.057*
C2	0.6104 (7)	0.5245 (5)	0.4445 (4)	0.062 (2)
H2A	0.6567	0.5813	0.4609	0.075*
H2B	0.5212	0.5426	0.4290	0.075*
C3	0.6020 (7)	0.4566 (5)	0.5099 (4)	0.062 (2)
H3A	0.5491	0.4844	0.5505	0.074*
H3B	0.6906	0.4446	0.5297	0.074*
C4	0.5388 (6)	0.3628 (5)	0.4847 (3)	0.0456 (16)
C5	0.6284 (6)	0.3133 (4)	0.4264 (3)	0.0413 (15)
H5A	0.7135	0.2947	0.4495	0.050*
C6	0.5531 (6)	0.2269 (4)	0.3963 (3)	0.0416 (15)
H6A	0.4708	0.2207	0.4259	0.050*
C7	0.5146 (6)	0.2506 (4)	0.3142 (3)	0.0385 (14)
H7A	0.4242	0.2279	0.3037	0.046*
C8	0.6122 (6)	0.2102 (4)	0.2542 (3)	0.0420 (15)
C9	0.7520 (6)	0.2353 (4)	0.2812 (3)	0.0417 (15)
H9A	0.7690	0.1973	0.3271	0.050*
C10	0.7671 (6)	0.3403 (4)	0.3048 (3)	0.0436 (16)
H10A	0.8465	0.3426	0.3373	0.052*
C11	0.6509 (6)	0.3782 (4)	0.3553 (3)	0.0373 (14)
C12	0.8015 (8)	0.3998 (5)	0.2341 (4)	0.068 (2)
H12A	0.7441	0.4551	0.2330	0.082*
H12B	0.8926	0.4220	0.2387	0.082*
C13	0.7873 (7)	0.3479 (6)	0.1600 (4)	0.067 (2)
H13A	0.8244	0.3890	0.1202	0.080*
C14	0.8569 (7)	0.2086 (5)	0.2239 (4)	0.0540 (18)
C15	0.5827 (7)	0.2387 (5)	0.1723 (4)	0.0566 (19)
H15A	0.4868	0.2451	0.1678	0.068*
H15B	0.6085	0.1859	0.1400	0.068*
C16	0.6447 (7)	0.3279 (5)	0.1382 (4)	0.070 (2)
H16A	0.6399	0.3232	0.0830	0.084*
H16B	0.5909	0.3819	0.1533	0.084*
C17	0.5155 (5)	0.3605 (4)	0.3181 (3)	0.0377 (14)
H17A	0.5149	0.3869	0.2664	0.045*
C18	0.5181 (7)	0.3020 (5)	0.5555 (3)	0.066 (2)
H18C	0.6006	0.2968	0.5829	0.098*
H18D	0.4886	0.2400	0.5406	0.098*
H18E	0.4522	0.3309	0.5877	0.098*
C19	0.4026 (6)	0.3801 (5)	0.4445 (3)	0.0461 (16)
H19A	0.3485	0.3235	0.4504	0.055*
H19B	0.3572	0.4313	0.4708	0.055*
C20	0.2808 (6)	0.3924 (5)	0.3252 (4)	0.061 (2)
H20A	0.2541	0.3264	0.3277	0.073*
H20B	0.2904	0.4091	0.2718	0.073*
C21	0.1728 (7)	0.4528 (6)	0.3602 (5)	0.081 (3)
H21A	0.0972	0.4544	0.3267	0.122*

H21B	0.2057	0.5161	0.3676	0.122*
H21C	0.1470	0.4265	0.4085	0.122*
C22	0.7553 (7)	0.6160 (5)	0.3047 (5)	0.082 (3)
H22A	0.7518	0.6418	0.2540	0.123*
H22B	0.8425	0.5905	0.3140	0.123*
H22C	0.7368	0.6651	0.3411	0.123*
C23	0.5638 (7)	0.0621 (5)	0.4222 (4)	0.0532 (18)
C24	0.6515 (7)	−0.0215 (4)	0.4266 (4)	0.0470 (16)
C25	0.6195 (8)	−0.0936 (5)	0.4745 (4)	0.065 (2)
H25A	0.5420	−0.0897	0.5034	0.078*
C26	0.7004 (10)	−0.1734 (6)	0.4811 (5)	0.083 (3)
H26A	0.6796	−0.2213	0.5156	0.100*
C27	0.8113 (10)	−0.1796 (5)	0.4357 (5)	0.082 (3)
H27A	0.8664	−0.2325	0.4388	0.099*
C28	0.8405 (8)	−0.1089 (5)	0.3865 (5)	0.073 (2)
H28A	0.9148	−0.1147	0.3551	0.087*
C29	0.7625 (7)	−0.0279 (5)	0.3814 (4)	0.0565 (18)
H29A	0.7851	0.0209	0.3481	0.068*

Atomic displacement parameters (Å^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.046 (3)	0.045 (2)	0.077 (3)	−0.004 (2)	−0.001 (3)	0.009 (2)
O2	0.040 (2)	0.036 (2)	0.050 (3)	0.002 (2)	0.005 (2)	0.008 (2)
O3	0.057 (3)	0.043 (2)	0.056 (3)	0.003 (2)	−0.005 (2)	−0.008 (2)
O4	0.062 (3)	0.075 (3)	0.053 (3)	−0.001 (3)	0.014 (3)	−0.005 (3)
O5	0.058 (3)	0.093 (4)	0.088 (4)	0.023 (3)	0.005 (3)	−0.024 (4)
O6	0.051 (3)	0.059 (3)	0.187 (6)	−0.004 (3)	0.018 (4)	0.014 (4)
N1	0.039 (3)	0.049 (3)	0.043 (3)	0.009 (3)	0.000 (3)	−0.003 (3)
C1	0.038 (3)	0.043 (4)	0.061 (4)	−0.003 (3)	−0.005 (3)	0.002 (4)
C2	0.058 (4)	0.049 (4)	0.080 (5)	0.009 (4)	−0.005 (4)	−0.023 (4)
C3	0.063 (5)	0.072 (5)	0.050 (4)	0.011 (4)	−0.003 (4)	−0.022 (4)
C4	0.039 (3)	0.057 (4)	0.041 (4)	0.013 (3)	0.002 (3)	−0.007 (3)
C5	0.036 (3)	0.044 (3)	0.044 (4)	0.004 (3)	0.003 (3)	0.001 (3)
C6	0.036 (3)	0.043 (4)	0.046 (4)	0.006 (3)	0.004 (3)	0.000 (3)
C7	0.038 (3)	0.038 (3)	0.040 (3)	−0.004 (3)	−0.003 (3)	0.000 (3)
C8	0.045 (4)	0.037 (3)	0.044 (4)	0.002 (3)	0.000 (3)	−0.005 (3)
C9	0.042 (3)	0.047 (4)	0.036 (3)	0.003 (3)	0.001 (3)	−0.008 (3)
C10	0.033 (3)	0.054 (4)	0.044 (4)	−0.003 (3)	−0.002 (3)	0.003 (3)
C11	0.033 (3)	0.039 (3)	0.040 (3)	0.003 (3)	0.003 (3)	−0.003 (3)
C12	0.078 (5)	0.068 (5)	0.058 (5)	−0.013 (4)	0.023 (4)	0.001 (4)
C13	0.066 (5)	0.082 (6)	0.053 (5)	−0.005 (5)	0.011 (4)	0.012 (4)
C14	0.038 (4)	0.062 (5)	0.062 (5)	0.002 (4)	−0.005 (4)	−0.013 (4)
C15	0.056 (4)	0.062 (4)	0.052 (4)	0.002 (4)	−0.007 (4)	0.005 (4)
C16	0.066 (5)	0.093 (6)	0.050 (4)	0.000 (5)	0.005 (4)	0.006 (4)
C17	0.033 (3)	0.045 (3)	0.036 (3)	0.003 (3)	0.001 (3)	0.000 (3)
C18	0.070 (5)	0.089 (6)	0.038 (4)	0.010 (5)	0.011 (4)	0.002 (4)
C19	0.040 (3)	0.049 (4)	0.049 (4)	0.006 (3)	0.007 (3)	0.002 (3)

C20	0.041 (4)	0.087 (6)	0.053 (4)	0.012 (4)	−0.005 (4)	0.000 (4)
C21	0.053 (5)	0.094 (6)	0.097 (7)	0.024 (5)	0.001 (5)	0.004 (6)
C22	0.061 (5)	0.050 (4)	0.136 (8)	−0.014 (4)	0.012 (6)	0.010 (5)
C23	0.049 (4)	0.046 (4)	0.064 (5)	−0.006 (4)	−0.002 (4)	0.006 (4)
C24	0.056 (4)	0.036 (3)	0.050 (4)	−0.006 (3)	−0.009 (4)	0.005 (3)
C25	0.076 (5)	0.051 (4)	0.068 (5)	−0.004 (5)	0.004 (5)	0.004 (4)
C26	0.106 (7)	0.063 (6)	0.081 (6)	0.002 (5)	−0.017 (6)	0.021 (5)
C27	0.108 (7)	0.041 (4)	0.099 (7)	0.023 (5)	−0.042 (6)	0.003 (5)
C28	0.073 (5)	0.066 (5)	0.079 (5)	0.028 (5)	−0.021 (5)	−0.017 (5)
C29	0.068 (5)	0.043 (4)	0.059 (4)	−0.003 (4)	−0.013 (4)	−0.004 (4)

Geometric parameters (Å, °)

O1—C22	1.416 (7)	C10—H10A	0.9800
O1—C1	1.443 (7)	C11—C17	1.528 (8)
O2—C23	1.325 (7)	C12—C13	1.496 (9)
O2—C6	1.454 (7)	C12—H12A	0.9700
O3—C8	1.434 (7)	C12—H12B	0.9700
O3—H3	0.8200	C13—C16	1.509 (10)
O4—C14	1.339 (8)	C13—H13A	0.9800
O4—C13	1.458 (9)	C15—C16	1.527 (9)
O5—C14	1.213 (8)	C15—H15A	0.9700
O6—C23	1.194 (8)	C15—H15B	0.9700
N1—C17	1.457 (7)	C16—H16A	0.9700
N1—C20	1.463 (7)	C16—H16B	0.9700
N1—C19	1.463 (7)	C17—H17A	0.9800
C1—C2	1.518 (8)	C18—H18C	0.9600
C1—C11	1.544 (8)	C18—H18D	0.9600
C1—H1A	0.9800	C18—H18E	0.9600
C2—C3	1.495 (9)	C19—H19A	0.9700
C2—H2A	0.9700	C19—H19B	0.9700
C2—H2B	0.9700	C20—C21	1.509 (9)
C3—C4	1.532 (9)	C20—H20A	0.9700
C3—H3A	0.9700	C20—H20B	0.9700
C3—H3B	0.9700	C21—H21A	0.9600
C4—C18	1.523 (8)	C21—H21B	0.9600
C4—C5	1.529 (8)	C21—H21C	0.9600
C4—C19	1.556 (8)	C22—H22A	0.9600
C5—C6	1.529 (8)	C22—H22B	0.9600
C5—C11	1.562 (8)	C22—H22C	0.9600
C5—H5A	0.9800	C23—C24	1.473 (9)
C6—C7	1.526 (8)	C24—C25	1.357 (9)
C6—H6A	0.9800	C24—C29	1.370 (9)
C7—C8	1.546 (8)	C25—C26	1.392 (10)
C7—C17	1.552 (8)	C25—H25A	0.9300
C7—H7A	0.9800	C26—C27	1.372 (12)
C8—C15	1.518 (8)	C26—H26A	0.9300
C8—C9	1.523 (8)	C27—C28	1.351 (10)
C9—C14	1.504 (8)	C27—H27A	0.9300

C9—C10	1.545 (8)	C28—C29	1.388 (9)
C9—H9A	0.9800	C28—H28A	0.9300
C10—C12	1.536 (8)	C29—H29A	0.9300
C10—C11	1.559 (8)		
C22—O1—C1	113.0 (5)	O4—C13—C12	110.3 (6)
C23—O2—C6	117.1 (4)	O4—C13—C16	111.7 (6)
C8—O3—H3	109.5	C12—C13—C16	113.7 (7)
C14—O4—C13	115.6 (5)	O4—C13—H13A	106.9
C17—N1—C20	110.7 (5)	C12—C13—H13A	106.9
C17—N1—C19	117.9 (5)	C16—C13—H13A	106.9
C20—N1—C19	112.1 (5)	O5—C14—O4	119.0 (7)
O1—C1—C2	107.5 (5)	O5—C14—C9	123.2 (7)
O1—C1—C11	110.0 (5)	O4—C14—C9	117.7 (6)
C2—C1—C11	117.6 (5)	C8—C15—C16	120.6 (6)
O1—C1—H1A	107.1	C8—C15—H15A	107.2
C2—C1—H1A	107.1	C16—C15—H15A	107.2
C11—C1—H1A	107.1	C8—C15—H15B	107.2
C3—C2—C1	111.9 (5)	C16—C15—H15B	107.2
C3—C2—H2A	109.2	H15A—C15—H15B	106.8
C1—C2—H2A	109.2	C13—C16—C15	116.3 (6)
C3—C2—H2B	109.2	C13—C16—H16A	108.2
C1—C2—H2B	109.2	C15—C16—H16A	108.2
H2A—C2—H2B	107.9	C13—C16—H16B	108.2
C2—C3—C4	110.9 (5)	C15—C16—H16B	108.2
C2—C3—H3A	109.5	H16A—C16—H16B	107.4
C4—C3—H3A	109.5	N1—C17—C11	110.5 (4)
C2—C3—H3B	109.5	N1—C17—C7	115.8 (5)
C4—C3—H3B	109.5	C11—C17—C7	100.8 (5)
H3A—C3—H3B	108.1	N1—C17—H17A	109.8
C18—C4—C5	111.5 (5)	C11—C17—H17A	109.8
C18—C4—C3	107.9 (5)	C7—C17—H17A	109.8
C5—C4—C3	110.0 (5)	C4—C18—H18C	109.5
C18—C4—C19	109.7 (5)	C4—C18—H18D	109.5
C5—C4—C19	106.7 (5)	H18C—C18—H18D	109.5
C3—C4—C19	111.0 (5)	C4—C18—H18E	109.5
C4—C5—C6	107.7 (5)	H18C—C18—H18E	109.5
C4—C5—C11	110.4 (5)	H18D—C18—H18E	109.5
C6—C5—C11	105.3 (5)	N1—C19—C4	115.6 (5)
C4—C5—H5A	111.1	N1—C19—H19A	108.4
C6—C5—H5A	111.1	C4—C19—H19A	108.4
C11—C5—H5A	111.1	N1—C19—H19B	108.4
O2—C6—C7	116.6 (5)	C4—C19—H19B	108.4
O2—C6—C5	110.6 (5)	H19A—C19—H19B	107.4
C7—C6—C5	106.0 (5)	N1—C20—C21	112.9 (6)
O2—C6—H6A	107.8	N1—C20—H20A	109.0
C7—C6—H6A	107.8	C21—C20—H20A	109.0
C5—C6—H6A	107.8	N1—C20—H20B	109.0
C6—C7—C8	113.6 (5)	C21—C20—H20B	109.0
C6—C7—C17	100.1 (5)	H20A—C20—H20B	107.8

C8—C7—C17	113.3 (5)	C20—C21—H21A	109.5
C6—C7—H7A	109.8	C20—C21—H21B	109.5
C8—C7—H7A	109.8	H21A—C21—H21B	109.5
C17—C7—H7A	109.8	C20—C21—H21C	109.5
O3—C8—C15	106.8 (5)	H21A—C21—H21C	109.5
O3—C8—C9	105.6 (5)	H21B—C21—H21C	109.5
C15—C8—C9	114.3 (5)	O1—C22—H22A	109.5
O3—C8—C7	108.0 (5)	O1—C22—H22B	109.5
C15—C8—C7	114.9 (5)	H22A—C22—H22B	109.5
C9—C8—C7	106.6 (5)	O1—C22—H22C	109.5
C14—C9—C8	112.3 (5)	H22A—C22—H22C	109.5
C14—C9—C10	110.5 (5)	H22B—C22—H22C	109.5
C8—C9—C10	113.4 (5)	O6—C23—O2	123.9 (7)
C14—C9—H9A	106.7	O6—C23—C24	123.7 (7)
C8—C9—H9A	106.7	O2—C23—C24	112.3 (6)
C10—C9—H9A	106.7	C25—C24—C29	120.0 (7)
C12—C10—C9	109.3 (5)	C25—C24—C23	119.3 (7)
C12—C10—C11	116.0 (5)	C29—C24—C23	120.6 (6)
C9—C10—C11	114.0 (5)	C24—C25—C26	121.3 (8)
C12—C10—H10A	105.5	C24—C25—H25A	119.4
C9—C10—H10A	105.5	C26—C25—H25A	119.4
C11—C10—H10A	105.5	C27—C26—C25	118.5 (8)
C17—C11—C1	116.3 (5)	C27—C26—H26A	120.8
C17—C11—C10	111.6 (5)	C25—C26—H26A	120.8
C1—C11—C10	107.8 (5)	C28—C27—C26	119.9 (8)
C17—C11—C5	96.6 (5)	C28—C27—H27A	120.0
C1—C11—C5	113.2 (5)	C26—C27—H27A	120.0
C10—C11—C5	111.1 (5)	C27—C28—C29	121.8 (8)
C13—C12—C10	114.3 (6)	C27—C28—H28A	119.1
C13—C12—H12A	108.7	C29—C28—H28A	119.1
C10—C12—H12A	108.7	C24—C29—C28	118.4 (7)
C13—C12—H12B	108.7	C24—C29—H29A	120.8
C10—C12—H12B	108.7	C28—C29—H29A	120.8
H12A—C12—H12B	107.6		

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
O3—H3···O1^{i}	0.82	2.25	3.056 (8)	166

Symmetry codes: (i) −*x*+1, *y*−1/2, −*z*+1/2.

Fig. 1

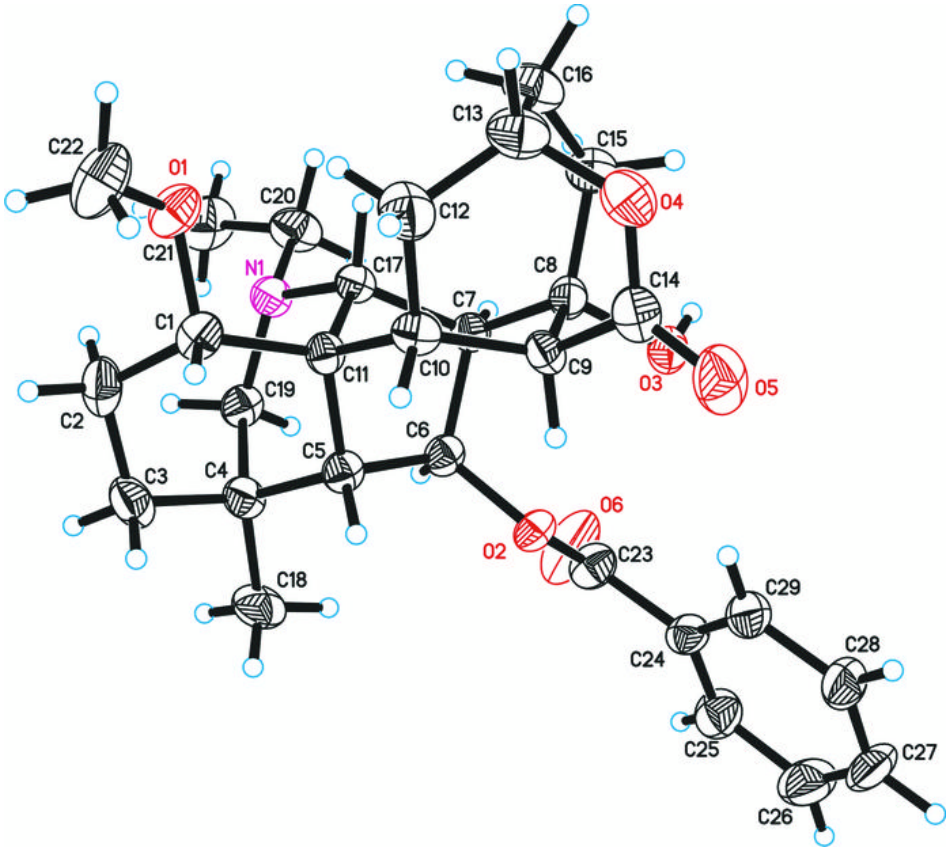
C16
C13
C15
C22
O1
C20
C12
O4
C21
N1
C17
C8
C14
C7
C1
C11
C10
C9
O3
O5
C19
C2
C6
C5
C3
C4
O6
O2
C23
C18
C29
C24
C28
C25
C27
C26

Fig. 2

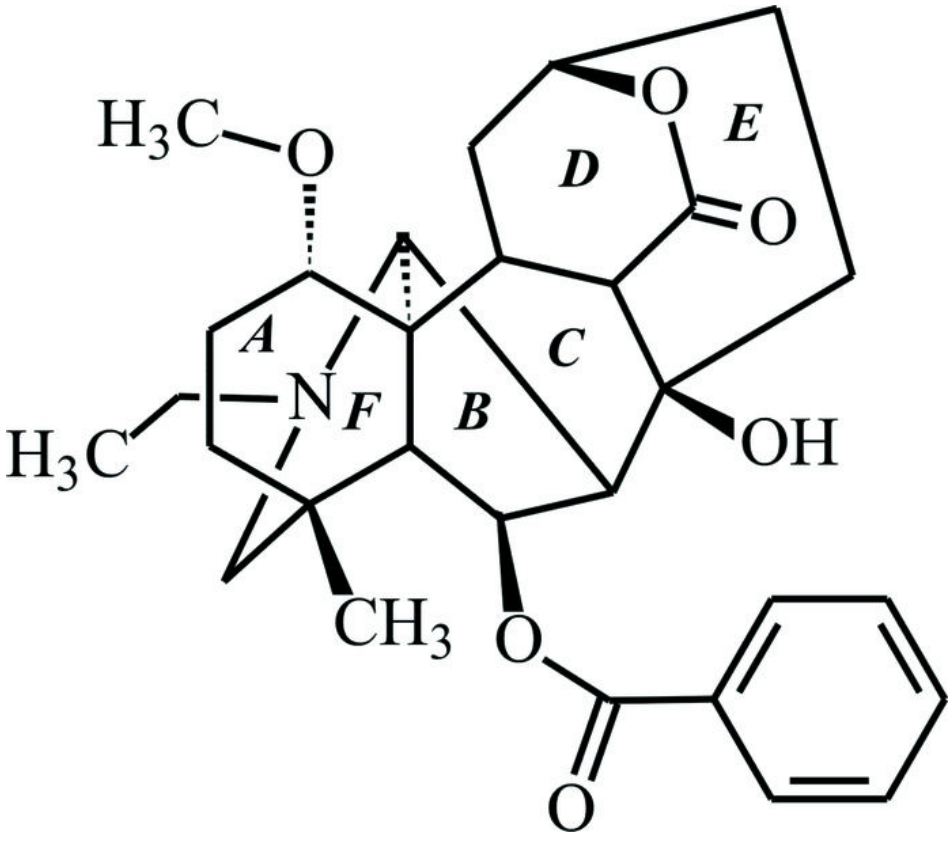

Fig. 3