

Berechnung von mittleren Schwingungsamplituden nach der Methode der charakteristischen Schwingungen von Müller

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(Z. Naturforsch. 25 a, 1292—1295 [1970]; eingegangen am 2. Mai 1970)

It is shown for many molecules and ions that the „Methode der charakteristischen Schwingungen“ of Müller is useful for the calculation of mean amplitudes of vibrations if the vibrations are rather characteristic.

In den letzten Jahren wurden sehr häufig mittlere Schwingungsamplituden aus spektroskopischen Daten berechnet. Da diese Rechnungen in vielen Fällen nicht einfach sind, wurden dabei oft fehlerhafte Ergebnisse erzielt.

Meistens wurden mittlere Schwingungsamplituden aus der Säkulargleichung:

$$|\sum \mathbf{G}^{-1} - \Delta \mathbf{E}| = 0 \quad (1)$$

berechnet¹. Es wurde bereits erwähnt, daß es sinnvoller ist, die \sum -Matrix aus den Eigenvektoren nach:

$$\sum = \mathbf{L} \Delta \mathbf{L}' \quad (2)$$

zu berechnen^{2,3}. In den Gln. (1) und (2) bedeutet \sum die symmetrisierte mittlere-quadratische Amplitudenmatrix mit den Elementen $\sum_{ij} = \langle S_i S_j \rangle$, S_i und S_j sind Symmetriekoordinaten, \mathbf{G}^{-1} ist die Matrix der kinetischen Energie, \mathbf{E} die Einheitsmatrix, Δ eine Diagonalmatrix mit den Elementen:

$$\Delta_i = \frac{h}{8\pi^2 \nu_i} \cdot \coth \frac{h \nu_i}{2kT}$$

und \mathbf{L} , die Eigenvektormatrix, die die Transformationsmatrix zwischen inneren Symmetriekoordinaten \mathbf{S} und Normalkoordinaten \mathbf{Q} gemäß $\mathbf{S} = \mathbf{L} \mathbf{Q}$ darstellt.

Bekanntlich gilt zwischen der inversen Matrix der kinetischen Energie \mathbf{G} und der \mathbf{L} -Matrix die Beziehung

$$\mathbf{G} = \mathbf{L} \mathbf{L}' \quad (3)$$

Es wurde festgestellt, daß man in vielen Fällen die Näherung $L_{ij} = 0$ für $i < j$ einführen kann^{2,4,5}, was

zu erheblicher Vereinfachung des Rechenproblems führt. Die Berechnung der \sum -Matrix kann allgemein nach Gl. (2) erfolgen; die \mathbf{L} -Matrix wird nach Gl. (3) mit der oben eingeführten Näherung ermittelt^{4,5}.

MÜLLER und Mitarb.⁶ haben vor kurzem eine Näherungsmethode entwickelt, mit der es ohne Lösung eines inversen Eigenwertproblems möglich ist, mittlere Schwingungsamplituden im Falle des Vorliegens charakteristischer Frequenzen näherungsweise zu berechnen. Bei dieser Methode wird ein gegebenes kompliziertes Molekül in Bruchstücke von je drei Atome des Typs XY_2 bzw. XYZ zerlegt, und die Schwingungsfrequenzen dieser „Pseudo-Moleküle“ abgeschätzt. Mit diesen Werten lassen sich mittlere Schwingungsamplituden aus den \sum -Elementen für $n=2$ (XY_2 -Bruchstücke) und $n=3$ (XYZ -Bruchstücke), aus den einfachen bereits angegebenen Formeln⁶ berechnen.

In ihrer Arbeit haben MÜLLER und Mitarb.⁶ diese Methode an einigen Beispielen erläutert und gezeigt, daß die so erhaltenen Werte gut mit denjenigen, die mit einer vollständigeren Methode erhalten wurden, übereinstimmen. Es ist der Zweck dieser Arbeit, mittlere Schwingungsamplituden für eine größere Reihe von Molekülen nach der beschriebenen Näherung zu berechnen, um somit die Gültigkeit derselben an weiteren Beispielen zu prüfen und zu bestätigen.

Aus Tab. 1 sind die mittleren Schwingungsamplituden für eine Reihe von Molekülen und Ionen des Typs ZXY_3 (C_{3v}) zu entnehmen und aus Tab. 2 die

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¹ S. J. CYVIN, Mean Square Amplitudes and Molecular Vibrations, Elsevier, Amsterdam 1968.

² A. MÜLLER, Z. Phys. Chem. Leipzig **238**, 116 [1968].

³ A. MÜLLER, B. KREBS, A. FADINI, O. GLEMSE, S. J. CYVIN, J. BRUNVOLL, B. N. CYVIN, I. ELVEBREDD, G. HAGEN u. B. VIZI, Z. Naturforsch. **23 a**, 1656 [1968].

⁴ C. J. PEACOCK u. A. MÜLLER, J. Mol. Spectr. **26**, 454 [1968].

⁵ C. J. PEACOCK, U. HEIDBORN u. A. MÜLLER, J. Mol. Spectr. **30**, 338 [1969].

⁶ A. MÜLLER, C. J. PEACOCK, H. SCHULZE u. U. HEIDBORN, J. Mol. Struct. **3**, 252 [1969].

ZXY ₃	T(°K)	u(XY)	u(XZ)	u(Y...Y)	u(Y...Z)	ZXY ₃	T(°K)	u(XY)	u(XY)	u(Y...Y)	u(Y...Z)		
OPCl ₃	298,2	a	0,0464	0,0354	0,085	0,068	BrGeH ₃	298,2	a	0,0894	0,0477	0,156	0,134
		b	0,0470	—	0,079	—			b	0,0894	—	0,155	—
		c	0,0472	0,0359	—	0,069			c	0,0894	0,0477	—	0,145
	0	a	0,0428	0,0353	0,060	0,058		0	a	0,0894	0,0379	0,155	0,128
		b	0,0435	—	0,056	—			b	0,0894	—	0,154	—
		c	0,0437	0,0357	—	0,059			c	0,0894	0,0379	—	0,137
OPBr ₃	298,2	a	0,0464	0,0359	0,090	0,077	IGeH ₃	298,2	a	0,0895	0,0517	0,157	0,143
		b	0,0507	—	0,084	—			b	0,0895	—	0,156	—
		c	0,0490	0,0371	—	0,078			c	0,0895	0,0524	—	0,151
	0	a	0,0407	0,0357	0,051	0,059		0	a	0,0895	0,0380	0,155	0,134
		b	0,0435	—	0,050	—			b	0,0895	—	0,155	—
		c	0,0427	0,0366	—	0,061			c	0,0895	0,0384	—	0,141
HPO ₃ ²⁻	298,2	a	0,0395	0,0862	0,063	0,109	HGeCl ₃	298,2	a	0,0461	0,0886	0,102	0,137
		b	0,0395	—	0,060	—			b	0,0465	—	0,098	—
		c	0,0392	0,0863	—	0,108			c	0,0462	0,0887	—	0,131
	0	a	0,0392	0,0862	0,058	0,109		0	a	0,0405	0,0886	0,066	0,130
		b	0,0392	—	0,056	—			b	0,0408	—	0,064	—
		c	0,0389	0,0863	—	0,107			c	0,0406	0,0887	—	0,127
HOPO ₃ ²⁻	298,2	a	0,0398	0,0415	0,059	0,067	HGeBr ₃	298,2	a	0,0476	0,0895	0,109	0,142
		b	0,0397	—	0,059	—			b	0,0488	—	0,104	—
		c	0,0394	0,0420	—	0,069			c	0,0478	0,0899	—	0,133
	0	a	0,0394	0,0409	0,056	0,060		0	a	0,0376	0,0895	0,056	0,132
		b	0,0394	—	0,056	—			b	0,0384	—	0,055	—
		c	0,0391	0,0415	—	0,061			c	0,0380	0,0897	—	0,129
OVF ₃	298,2	a	0,0404	0,0366	0,098	0,079	ClGeBr ₃	298,2	a	0,0477	0,0472	0,111	0,099
		b	0,0409	—	0,092	—			b	0,0498	—	0,109	—
		c	0,0410	0,0365	—	0,076			c	0,0481	0,0465	—	0,098
	0	a	0,0394	0,0364	0,072	0,065		0	a	0,0376	0,0411	0,057	0,060
		b	0,0399	—	0,070	—			b	0,0387	—	0,057	—
		c	0,0399	0,0362	—	0,064			c	0,0381	0,0410	—	0,059
OVCl ₃	298,2	a	0,0459	0,0369	0,112	0,090	FGeH ₃	298,2	a	0,0893	0,0415	0,156	0,125
		b	0,0470	—	0,108	—			b	0,0893	—	0,154	—
		c	0,0469	0,0372	—	0,085			c	0,0893	0,0418	—	0,131
	0	a	0,0411	0,0366	0,068	0,067		0	a	0,0893	0,0401	0,154	0,122
		b	0,0420	—	0,067	—			b	0,0893	—	0,153	—
		c	0,0420	0,0363	—	0,065			c	0,0893	0,0403	—	0,127
OVBr ₃	298,2	a	0,0465	0,0371	0,119	0,097	ClGeH ₃	298,2	a	0,0893	0,0464	0,156	0,134
		b	0,0498	—	0,112	—			b	0,0893	—	0,155	—
		c	0,0491	0,0381	—	0,092			c	0,0892	0,0467	—	0,141
	0	a	0,0384	0,0368	0,058	0,067		0	a	0,0893	0,0407	0,154	0,128
		b	0,0401	—	0,057	—			b	0,0893	—	0,153	—
		c	0,0402	0,0374	—	0,066			c	0,0893	0,0409	—	0,134
ClReO ₃	298,2	a	0,0346	0,0550	0,068	0,091	ISiBr ₃	298,2	a	0,0463	0,0648	0,118	0,087
		b	0,0345	—	0,063	—			b	0,0342	—	0,088	—
		c	0,0346	0,0563	—	0,099			c	0,0339	0,0565	—	0,126
	0	a	0,0343	0,0432	0,060	0,067		a	0,0414	0,0487	0,057	0,049	
		b	0,0342	—	0,057	—			b	0,0308	—	0,050	—
		c	0,0342	0,0439	—	0,071			c	0,0306	0,0399	—	0,057
BrReO ₃	298,2	a	0,0346	0,0571	0,070	0,099	ClSiH ₃	298,2	a	0,0888	0,0461	0,151	0,128
		b	0,0345	—	0,069	—			b	0,0888	—	0,150	—
		c	0,0345	0,0593	—	0,108			c	0,0895	0,0474	—	0,134
	0	a	0,0343	0,0382	0,061	0,066		0	a	0,0888	0,0431	0,150	0,125
		b	0,0342	—	0,060	—			b	0,0888	—	0,149	—
		c	0,0342	0,0393	—	0,070			c	0,0895	0,0441	—	0,130
SReO ₃	298,2	a	0,0360	0,0384	0,074	0,077	BrSiH ₃	298,2	a	0,0888	0,0476	0,151	0,132
		b	0,0358	—	0,073	—			b	0,0888	—	0,150	—
		c	0,0360	0,0382	—	0,081			c	0,0888	0,0492	—	0,138
	0	a	0,0357	0,0351	0,063	0,062		0	a	0,0888	0,0422	0,150	0,127
		b	0,0354	—	0,062	—			b	0,0888	—	0,149	—
		c	0,0355	0,0350	—	0,063			c	0,0888	0,0434	—	0,132

Tab. 1

ZXY ₃	T(°K)	u(XY)	u(XZ)	u(Y...Y)	u(Y...Z)	ZXY ₃	T(°K)	u(XY)	u(XZ)	u(Y...Y)	u(Y...Z)		
ISiH ₃	298,2	a	0,0887	0,0516	0,152	0,139	SSO ₃ ²⁻	298,2	a	0,0390	0,0467	0,059	0,069
		b	0,0888	—	0,151	—			b	0,0389	—	0,056	—
		c	0,0887	0,0536	—	0,146			c	0,0386	0,0541	—	0,073
	0	a	0,0887	0,0435	0,151	0,132		0	a	0,0387	0,0429	0,056	0,059
		b	0,0888	—	0,150	—			b	0,0386	—	0,053	—
		c	0,0887	0,0449	—	0,138			c	0,0384	0,0482	—	0,062
HSiF ₃	298,2	a	0,0402	0,0865	0,075	0,117	HSO ₃ ²⁻	298,2	a	0,0375	0,0812	0,059	0,104
		b	0,0404	—	0,071	—			b	0,0376	—	0,057	—
		c	0,0399	0,0866	—	0,115			c	0,0372	0,0813	—	0,102
	0	a	0,0397	0,0865	0,063	0,115		0	a	0,0373	0,0812	0,055	0,103
		b	0,0399	—	0,061	—			b	0,0375	—	0,054	—
		c	0,0395	0,0866	—	0,113			c	0,0371	0,0813	—	0,102
HSiCl ₃	298,2	a	0,0464	0,0876	0,090	0,129	FSO ₃ ²⁻	298,2	a	0,0364	0,0426	0,055	0,064
		b	0,0480	—	0,086	—			b	0,0366	—	0,055	—
		c	0,0472	0,0897	—	0,122			c	0,0361	0,0433	—	0,065
	0	a	0,0432	0,0876	0,062	0,125		0	a	0,0363	0,0416	0,053	0,058
		b	0,0445	—	0,061	—			b	0,0365	—	0,053	—
		c	0,0441	0,0879	—	0,120			c	0,0360	0,0423	—	0,059
HSiBr ₃	298,2	a	0,0476	0,0880	0,094	0,134	HOSO ₃ ²⁻	298,2	a	0,0377	0,0411	0,056	0,063
		b	0,0502	—	0,091	—			b	0,0376	—	0,055	—
		c	0,0489	0,0887	—	0,124			c	0,0372	0,0419	—	0,064
	0	a	0,0420	0,0880	0,053	0,127		0	a	0,0374	0,0405	0,053	0,058
		b	0,0438	—	0,052	—			b	0,0374	—	0,053	—
		c	0,0432	0,0885	—	0,122			c	0,0371	0,0413	—	0,059
FSiCl ₃	298,2	a	0,0416	0,0413	0,092	0,078	HOSeO ₃ ²⁻	298,2	a	0,0381	0,0411	0,068	0,074
		b	0,0453	—	0,094	—			b	0,0382	—	0,067	—
		c	0,0406	0,0410	—	0,076			c	0,0380	0,0414	—	0,075
	0	a	0,0395	0,0407	0,062	0,062		0	a	0,0376	0,0400	0,061	0,064
		b	0,0422	—	0,064	—			b	0,0377	—	0,060	—
		c	0,0392	0,0405	—	0,061			c	0,0375	0,0403	—	0,064
BrSiCl ₃	298,2	a	0,0460	0,0502	0,106	0,077	FCrO ₃	298,2	a	0,0389	0,0454	0,071	0,086
		b	0,0457	—	0,087	—			b	0,0389	—	0,071	—
		c	0,0453	0,0551	—	0,109			c	0,0387	0,0457	—	0,088
	0	a	0,0429	0,0433	0,066	0,053		0	a	0,0384	0,0433	0,063	0,069
		b	0,0430	—	0,061	—			b	0,0384	—	0,062	—
		c	0,0427	0,0465	—	0,063			c	0,0383	0,0436	—	0,069
ClSiBr ₃	298,2	a	0,0477	0,0471	0,100	0,084	ClCrO ₃	298,2	a	0,0388	0,0485	0,072	0,093
		b	0,0513	—	0,080	—			b	0,0389	—	0,072	—
		c	0,0671	0,0477	—	0,087			c	0,0387	0,0481	—	0,099
	0	a	0,0419	0,0437	0,054	0,056		0	a	0,0384	0,0426	0,063	0,069
		b	0,0441	—	0,049	—			b	0,0385	—	0,063	—
		c	0,0525	0,0440	—	0,057			c	0,0383	0,0426	—	0,071
ISiCl ₃	298,2	a	0,0467	0,0510	0,115	0,078	NOsO ₃	298,2	a	0,0366	0,0359	0,069	0,076
		b	0,0467	—	0,109	—			b	0,0365	—	0,069	—
		c	0,0461	0,0567	—	0,115			c	0,0366	0,0358	—	0,076
	0	a	0,0435	0,0429	0,069	0,052		0	a	0,0360	0,0356	0,061	0,065
		b	0,0437	—	0,068	—			b	0,0360	—	0,061	—
		c	0,0433	0,0464	—	0,063			c	0,0360	0,0356	—	0,065

Tab. 1. Mittlere Schwingungsamplituden für ZXY₃(C_{3v})-Moleküle. Werte in Å, erhalten von: a) vollständige Rechnung³; b) XY₂-Bruchstück; c) ZXY-Bruchstück.

entsprechenden Werte für Moleküle des Typs ZXY₂ (C_{2v}). Die Rechnungen wurden für zwei Temperaturen (298,2 und 0 °K) durchgeführt und die erhaltenen Werte jeweils mit den Werten verglichen, die mit vollständiger Rechnung erhalten wurden. Die Schwingungsfrequenzen sind die gleichen wie bei³. Zur Mittelung bei entarteten Schwingungen vgl.⁶

Wie leicht zu ersehen ist, liefern die Näherungsrechnungen brauchbare Werte, vor allem wenn das Zentralatom relativ schwerer ist; aber auch in allen Fällen und auch bei den Amplituden für nicht gebundene Atome [$u(Y...Y)$ und $u(Y...Z)$] ist die Übereinstimmung relativ gut. Die Unterschiede zu den Literaturwerten sind lediglich in wenigen

ZXY ₂	T(°K)	u(XY)	u(XZ)	u(Y...Y)	u(Y...Z)	ZXY ₂	T(°K)	u(XY)	u(XZ)	u(Y...Y)	u(Y...Z)		
OCH ₂	298,2	a	0,0805	0,0377	0,121	0,096	SCCl ₂	298,2	a	0,0509	0,0420	0,083	0,065
		b	0,0804	—	0,120	—			b	0,0555	—	0,091	—
		c	0,0804	0,0374	—	0,094			c	0,0556	0,0380	—	0,051
	0	a	0,0805	0,0377	0,121	0,096		0	a	0,0491	0,0416	0,059	0,052
		b	0,0804	—	0,120	—			b	0,0527	—	0,063	—
		c	0,0804	0,0374	—	0,094			c	0,0533	0,0375	—	0,040
OCD ₂	298,2	a	0,0686	0,0375	0,100	0,083	FCIF ₂	298,2	a	0,0493	0,0450	0,085	0,064
		b	0,0682	—	0,100	—			b	0,0505	—	0,077	—
		c	0,0683	0,0379	—	0,079			c	0,0495	0,0473	—	0,067
	0	a	0,0686	0,0375	0,100	0,082		0	a	0,0466	0,0435	0,070	0,059
		b	0,0682	—	0,100	—			b	0,0476	—	0,066	—
		c	0,0683	0,0379	—	0,079			c	0,0471	0,0425	—	0,060
HCO ₂ ⁻	298,2	a	0,0409	0,0802	0,049	0,096	FBrF ₂	298,2	a	0,0468	0,0432	0,088	0,066
		b	0,0404	—	0,046	—			b	0,0473	—	0,086	—
		c	0,0409	0,0807	—	0,091			c	0,0471	0,0424	—	0,072
	0	a	0,0408	0,0802	0,048	0,096		0	a	0,0438	0,0412	0,071	0,059
		b	0,0404	—	0,046	—			b	0,0442	—	0,070	—
		c	0,0408	0,0806	—	0,091			c	0,0441	0,0407	—	0,063
DCO ₂ ⁻	298,2	a	0,0408	0,0683	0,049	0,083	HONO ₂	298,2	a	0,0390	0,0469	0,049	0,054
		b	0,0407	—	0,047	—			b	0,0385	—	0,050	—
		c	0,0412	0,0684	—	0,078			c	0,0390	0,0485	—	0,051
	0	a	0,0407	0,0683	0,048	0,083		0	a	0,0389	0,0463	0,048	0,052
		b	0,0407	—	0,046	—			b	0,0385	—	0,049	—
		c	0,0411	0,0684	—	0,078			c	0,0390	0,0479	—	0,049
OCF ₂	298,2	a	0,0436	0,0362	0,053	0,050	DONO ₂	298,2	a	0,0391	0,0463	0,048	0,055
		b	0,0454	—	0,054	—			b	0,0387	—	0,047	—
		c	0,0457	0,0364	—	0,052			c	0,0392	0,0477	—	0,053
	0	a	0,0434	0,0362	0,051	0,048		0	a	0,0391	0,0458	0,047	0,053
		b	0,0452	—	0,051	—			b	0,0386	—	0,047	—
		c	0,0466	0,0363	—	0,050			c	0,0392	0,0472	—	0,051
OCCl ₂	298,2	a	0,0498	0,0369	0,068	0,058	FNO ₂	298,2	a	0,0380	0,0481	0,055	0,059
		b	0,0531	—	0,069	—			b	0,0376	—	0,058	—
		c	0,0532	0,0384	—	0,062			c	0,0381	0,0498	—	0,053
	0	a	0,0480	0,0368	0,054	0,052		0	a	0,0379	0,0472	0,052	0,055
		b	0,0511	—	0,054	—			b	0,0376	—	0,053	—
		c	0,0515	0,0383	—	0,056			c	0,0381	0,0489	—	0,051
OCBr ₂	298,2	a	0,0506	0,0368	0,071	0,062	ClNO ₂	298,2	a	0,0388	0,0434	0,056	0,063
		b	0,0564	—	0,072	—			b	0,0384	—	0,062	—
		c	0,0554	0,0392	—	0,072			c	0,0390	0,0452	—	0,062
	0	a	0,0474	0,0367	0,046	0,052		0	a	0,0387	0,0425	0,053	0,055
		b	0,0521	—	0,046	—			b	0,0384	—	0,056	—
		c	0,0523	0,0388	—	0,060			c	0,0389	0,0443	—	0,054
SCF ₂	298,2	a	0,0457	0,0385	0,056	0,053							
		b	0,0486	—	0,058	—							
		c	0,0486	0,0381	—	0,059							
	0	a	0,0453	0,0383	0,053	0,049							
		b	0,0480	—	0,055	—							
		c	0,0482	0,0380	—	0,052							

Tab. 2. Mittlere Schwingungsamplitude für ZXY₂(C_{2v})-Moleküle. Werte in Å, erhalten von: a) vollständige Rechnung²; b) XY₂-Bruchstück; c) ZXY-Bruchstück.

Fällen größer als 5%, und zwar dann, wenn die Masse der Außenatome im Verhältnis zur Masse des Zentralatoms groß ist und vor allen Dingen, wenn mehrere schwere Außenatome vorhanden sind (vgl. z. B. ISiBr₃, OPBr₃, HSiBr₃ etc.).

Man kann also zusammenfassend und an Hand unserer Rechnungen sagen, daß diese Methode mit geringem Rechenaufwand zu guten Ergebnissen führt und vor allen Dingen für Überschlagsrechnungen

außerordentlich brauchbar ist. Die Methode kann man allgemein für ein beliebiges Molekül anwenden, vorausgesetzt, daß die Zuordnung ihrer Schwingungsfrequenzen bekannt ist und charakteristische Schwingungen vorliegen.

Diese Arbeit wurde mit Unterstützung des „Consejo Nacional de Investigaciones Científicas y Técnicas de la República Argentina“ und einem Kontrakt der UNLP durchgeführt.

Fine Structure of the NH Stretching Band of Pyrrole

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(Z. Naturforsch. 25 a, 1296—1299 [1970]; received 2 May 1970)

The rotational fine structure of the fundamental NH stretching band of pyrrole has been analyzed. The analysis has been based on the symmetric top approximation. The obtained rotational constants $B''=0.3032\text{ cm}^{-1}$, $B'=0.3027\text{ cm}^{-1}$, and $C=0.1505\text{ cm}^{-1}$ are in good agreement with the microwave data. Also the accurate band center $\nu=3530.520\text{ cm}^{-1}$ has been obtained.

According to the microwave structural analysis of the pyrrole molecule^{1,2} the ratio of the two moments of inertia about the two perpendicular axes in the molecular plane is 0.986. Accordingly, the molecule is very close to an oblate symmetric top and the analysis of the rotational fine structure of the vibrational bands can be performed by applying the theory of the symmetric top molecules. The complete asymmetric top analysis of the infrared spectrum of a heavy molecule like pyrrole would be rather cumbersome and even impossible.

In this paper the fundamental NH stretching band of pyrrole has been analyzed because it fits the region of the high resolution spectrometer constructed in our laboratory. The obtained rotational constants are good approximations to the actual ones and may be compared with the microwave rotational constants. Further, the accurate band center has been obtained which is useful for the research of vibrational anharmonicities in progress at this laboratory.

Experimental

The high resolution spectrum of the NH stretching band of pyrrole was recorded by means of a large near-infrared Pfund-type grating spectrometer which has been described elsewhere³. The vapor sample was in a 3-meter multiple pass absorption cell in a pressure of about 5 mm Hg. The relative humidity inside the spectrometer was about 13 per cent causing the superimposed water vapor absorptions which could be identified with the help of the paper of PLYLER and TIDWELL⁴. The spectral slitwidth during the run was 0.14 cm^{-1} . The wavenumbers of the absorption lines were calibrated against the known emission lines of

rare gases. The pyrrole used in this work was purchased from Fluka AG and was of purissimum grade and used without any further purification.

Theoretical

The rotational term value which is applicable for the non-degenerate vibrational energy states is given by

$$F(J, K) = BJ(J+1) + (C-B)K^2 - D_J J^2(J+1)^2,$$

when the negligible centrifugal distortion terms dependent on K have been omitted. In this case B is the mean value of the actual rotational constants A and B of a pyrrole molecule.

The vibration of the NH stretching mode ν_9 of pyrrole is perpendicular to the top axis. According to the selection rules of a perpendicular band, $\Delta J = \pm 1$ and $\Delta K = \pm 1$, the wavenumbers of the sub band components can be expressed in terms of m which is $-J$ in the P branch and $J+1$ in the R branch of the band,

$$\nu = \nu_K^{\text{sub}} + (B' + B'')m + (B' - B'')m^2 - 4D_J m^3.$$

The sub band origins ν_K^{sub} are given by

$$\nu_K^{\text{sub}} = \nu_0 + (C' - B') \pm 2(C' - B')K + [(C' - B') - (C'' - B'')]K^2,$$

where ν_0 is the wavenumber of the band center. The $+$ and $-$ signs refer to the positive ($\Delta K = 1$) and negative ($\Delta K = -1$) sub bands, respectively. The rotational constants of the upper state are denoted by the single prime and those of the lower state by the double prime.

Sonderdruckanforderungen an Dr. J. ELORANTA, Laboratory of Molecular Spectroscopy, Department of Physics, University of Oulu, Kontinkangas, Oulu, Finland.

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Results

For the infrared fundamental bands the difference between the rotational constants of the upper and the lower state is small as a rule and thus the last term in the expression of the sub band origin should be small. The spacing of the subsequent sub bands should be $2(C' - B')$, which is about 0.3 cm^{-1} according to the microwave data. On the other hand the spacing of the J components should be

$$B' + B'' \sim 0.6 \text{ cm}^{-1}.$$

Accordingly, the sub band centers and the J components of different sub bands will be superimposed with the same interval of 0.3 cm^{-1} . One can expect the absorptions to coincide up to rather high values of J and K due to the planarity and the accidental symmetry of the molecule.

On the basis of the foregoing discussion of the structure of the band a theoretical band shape was computed using the Hönl-London intensity formulae⁵. The rotational constants reported by BAK et al.¹ and the temperature of 300°K were used. The difference between the primed rotational con-

stants was omitted. Fig. 1 illustrates the sub bands of the lowest five K transitions and the resultant of sub bands up to $K = 80$ when every other line has been left out. The small circles and triangles denote the portion of the negative and the positive sub bands, respectively. The cross lines indicate the portion of the Q lines in the pattern.

The calculated spectrum was employed as a reference in the assignment of the spectral lines of the experimental spectrum. Also the experimental spectrum is illustrated in Fig. 1 and one can see the good agreement with the calculated one.

The B constants should be calculated from single sub bands but because of the superposition of the sub bands the individual bands cannot be resolved. Each absorption line is a sum of the sub band lines of different m values and as can be seen in the computed spectrum the portion of the positive sub bands in the P branch is quite small as is the portion of the negative sub bands in the R branch. The actual asymmetry and on the other hand the non-zero coefficients $B' - B''$ and $4D$ may break the coincidence of the lines when m is high. Therefore it is

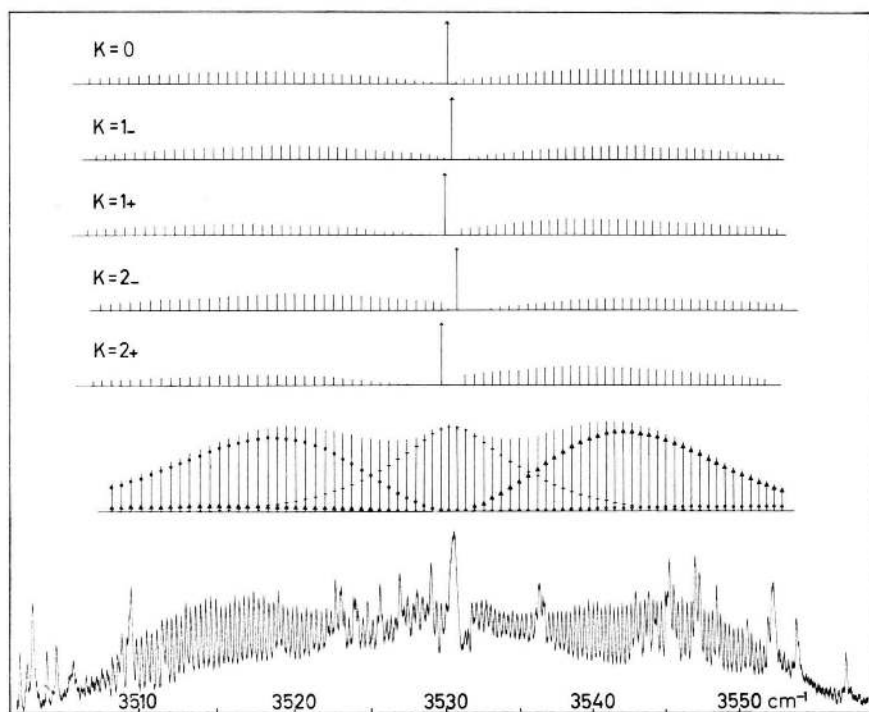


Fig. 1. Fine structure of the NH stretching band of pyrrole. The upper five spectra are the perpendicular type sub bands of the lowest K values computed with the Hönl-London intensity formulae at 300°K . The sixth spectrum is the sum of the sub bands. This spectrum is not to the same scale as the upper ones and every other line has been omitted for clarity. The circles and the triangles denote the portion of the negative and the positive sub bands in the line intensity, respectively. The portion of the Q components is indicated by the cross lines. The lowest spectrum is the recorded one. Except for the strong central maximum near 3531 cm^{-1} , which is the parallel part of the hybrid band, all the lines which differ from the calculated ones are due to water vapor absorptions.

⁵ G. HERZBERG, *Molecular Spectra and Molecular Structure, II. Infrared and Raman Spectra of Polyatomic Molecules*, D. van Nostrand Co., New York 1964, pp. 424-428.

not probable that a measured absorption maximum might present the wavenumber of the weak lines.

An attempt was made to compute the rotational constants separately for the positive sub bands from the R branch absorptions and for the negative sub bands from the P branch absorptions. The obtained results differed too greatly, probably due to the uneven distribution of the measurable lines over the entire band (cf. Table 1). Therefore, the least square fit over the whole band was used to evaluate the rotational constants. The fits were computed for the sub bands $K=0$, $K=1_-$, and $K=1_+$ simply by changing the numbering of the absorptions. In order to avoid unrealistic contribution from the unresolved sub band components, C' was taken as equal to C'' . The C constant as well as the band origin ν_0 were computed from the obtained sub band origins.

The observed and calculated wavenumbers and their differences are collected in Table 1. The calculated values have been taken from the polynomial computed for the sub band $K=0$,

$$\nu = 3530.375 + 0.60594 m - 5.10 \times 10^{-4} m^2 - 1 \times 10^{-6} m^3.$$

m	$\nu_{\text{obs.}}$ cm ⁻¹	$\nu_{\text{calc.}}$ cm ⁻¹	$\nu_{\text{obs.}} - \nu_{\text{calc.}}$ cm ⁻¹	m	$\nu_{\text{obs.}}$ cm ⁻¹	$\nu_{\text{calc.}}$ cm ⁻¹	$\nu_{\text{obs.}} - \nu_{\text{calc.}}$ cm ⁻¹
-30	3511.747	3511.766	-0.019	-15.5	3520.877	3520.865	0.012
-29.5	12.063	12.082	-0.019	-15	21.173	21.175	-0.002
-29	12.374	12.399	-0.025	-14.5	21.472	21.485	-0.013
-28.5	12.713	12.715	-0.002	-14	21.774	21.795	-0.021
-28	13.043	13.032	0.011	-12.5	22.690	22.723	-0.033
-27.5	13.362	13.348	0.014	12	37.558	37.571	-0.013
-27	13.669	13.663	0.006	13	38.168	38.164	0.004
-26.5	13.982	13.979	0.003	13.5	38.471	38.460	0.011
-26	14.305	14.294	0.011	14	38.764	38.756	0.008
-25.5	14.612	14.609	0.003	14.5	39.057	39.051	0.006
-25	14.917	14.924	-0.007	15	39.341	39.346	-0.005
-24.5	15.229	15.239	-0.010	15.5	39.628	39.641	-0.013
-24	15.556	15.553	0.003	16	39.912	39.936	-0.024
-23.5	15.866	15.867	-0.001	16.5	40.206	40.230	-0.024
-23	16.186	16.181	0.005	17	40.507	40.524	-0.017
-22.5	16.517	16.495	0.022	17.5	40.804	40.818	-0.014
-22	16.834	16.809	0.025	18	41.117	41.111	0.006
-21.5	17.141	17.122	0.019	18.5	41.422	41.404	0.018
-21	17.451	17.435	0.016	19	41.713	41.697	0.016
-20.5	17.768	17.748	0.020	19.5	42.009	41.990	0.019
-20	18.068	18.061	0.007	20	42.305	42.282	0.023
-19.5	18.367	18.373	-0.006	26.5	46.069	46.056	0.013
-19	18.675	18.685	-0.010	27	46.361	46.344	0.017
-18.5	18.966	18.997	-0.031	29	47.482	47.494	-0.012
-18	19.295	19.309	-0.014	29.5	47.785	47.781	0.004
-17.5	19.600	19.621	-0.021	30	48.086	48.067	0.019
-17	19.948	19.932	0.016	31.5	48.906	48.924	-0.018
-16.5	20.259	20.243	0.016	32	49.198	49.210	-0.012
-16	20.577	20.554	0.023	32.5	49.484	49.495	-0.011

Table 1. Observed and calculated wavenumbers of the fine structure components of the NH stretching band of pyrrole.

The half integers of m which appear in Table 1 refer to the absorptions which belong to the sub bands with odd K values. The average difference between the measured and calculated wavenumbers is 0.014 cm⁻¹.

The calculated rotational constants and the wavenumber of the band center are presented in Table 2 together with the microwave data. The coefficient of the third power term in the polynomial fits of various sub bands varied so much that the D_J constant could not be evaluated reliably.

This work	NYGAARD et al. ²	
$\nu_0 = 3530.520$		Table 2.
$B'' = 0.3032$	$A = 0.3046$	Molecular constants
$B' = 0.3027$	$B = 0.3003$	of pyrrole. All the
$C = 0.1505$	$C = 0.1512$	values are in cm ⁻¹ .

Discussion

The rotational constants obtained agree well with the microwave data ^{1, 2}. The differences are within the limits of the experimental errors. The difference

between B'' and B' is small, but it has been proved to be positive by a more appropriate graphical analysis.

Except for the center of the Q branch, the recorded absorption band agrees well with the computed one. The slight deviation from the exact symmetric top causes the greatest discrepancies when K is

© M. C. ALLEN and P. C. CROSS, *Molecular Vib-Rotors*, John Wiley & Sons Inc., New York 1963, p. 178.

small, because it is there that the strongest components of the parallel type band exist, and these transitions are allowed for in an asymmetric top molecule^{5, 6}.

The author wishes to thank Dr. R. ANTILA for great help in recording the spectra and Prof. P. TUOMIKOSKI for suggesting the topic. This work was financially supported by Neste Oy and the National Research Council for Sciences which are gratefully acknowledged.

Radiation-Induced Conformation Changes of the Pyrimidine Ring in a Single Crystal of Dihydrothymine

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(Z. Naturforsch. 25 a, 1299—1302 [1970]; received 10 January 1970)

Radiation-induced radicals in a single crystal of dihydrothymine were studied with EPR. It was found that the 5-thymyl radical is the dominant radical species at room temperature. The orientation dependence of the hyperfine couplings can only be explained if one assumes that $C_{(5)}$, $C_{(6)}$ and $C_{(7)}$ lie in the plane defined by the rest of the nonhydrogen atoms of the ring. The highly disordered structure of the ring in the undamaged crystal is ordered if a proton bound to $C_{(5)}$ is removed.

Radiation damage of several dihydropyrimidines in powder and solution has been studied with electron paramagnetic resonance (EPR)¹. No similar investigation in single crystals of any of these compounds has been done. In the present paper we report an EPR study of the structure of free radicals in a gamma-irradiated single crystal of dihydrothymine. The crystal and molecular structure of this compound has been reported by FURBERG and JENSEN². In contrast to thymine^{3, 4}, the pyrimidine ring in the crystal of dihydrothymine is highly disordered. From the EPR study of the radiation induced radicals in a single crystal of dihydrothymine it is possible to determine whether or not the disordered structure of the ring remains unchanged after the molecule has been damaged by irradiation.

Experimental

The crystals were grown by slow evaporation of an aqueous solution at 20 °C. No good quality crystals of moderate size could be found. Fortunately, dihydrothymine is very sensitive to irradiation and the crystal $1 \times 1 \times 0.2$ mm³ in size was sufficient for the EPR ana-

lysis. The external configuration and the crystallographic axes of the crystal used in this study are shown in Fig. 1. The axes were identified with X-ray diffraction analysis. The crystal is orthorhombic and the dimensions of the unit cell are $a=7.336$, $b=23.474$ and $c=7.034$ Å (l. c.²). There are eight molecules in the unit cell.

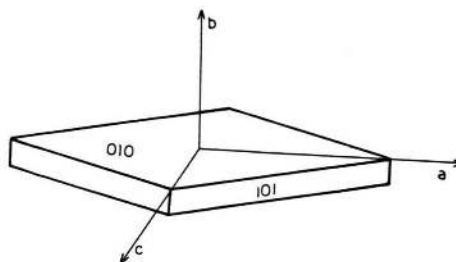


Fig. 1. External configuration and the crystallographic axes of the single crystal of dihydrothymine.

The crystal was irradiated in air at room temperature to a dose of about 5 Mrads by ⁶⁰Co gamma rays, at a dose rate of 0.3 Mrads/hour. The crystal was aligned in the EPR cavity so that one of the crystallographic axes was perpendicular to the magnetic field. The spectra were recorded at room temperature with a

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