



LUND UNIVERSITY

Lifetime measurements of the E (1)Pi, $\epsilon=0$ and $\epsilon=1$ states of (CO)-C-12-O-16, (CO)-C-13-O-16, and (CO)-C-13-O-18

Cacciani, P; Lynga, C; L'Huillier, Anne; Wahlström, Claes-Göran; Ubachs, W; Hinnen, P. C

Published in:
Astrophysical Journal

DOI:
[10.1086/311378](https://doi.org/10.1086/311378)

1998

[Link to publication](#)

Citation for published version (APA):
Cacciani, P., Lynga, C., L'Huillier, A., Wahlström, C.-G., Ubachs, W., & Hinnen, P. C. (1998). Lifetime measurements of the E (1)Pi, $\epsilon=0$ and $\epsilon=1$ states of (CO)-C-12-O-16, (CO)-C-13-O-16, and (CO)-C-13-O-18. *Astrophysical Journal*, 499. <https://doi.org/10.1086/311378>

Total number of authors:
6

General rights

Unless other specific re-use rights are stated the following general rights apply:
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

Read more about Creative commons licenses: <https://creativecommons.org/licenses/>

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

LUND UNIVERSITY

PO Box 117
221 00 Lund
+46 46-222 00 00

LIFETIME MEASUREMENTS OF THE $E\ ^1\Pi$, $v = 0$ AND $v = 1$ STATES OF $^{12}\text{C}^{16}\text{O}$, $^{13}\text{C}^{16}\text{O}$, AND $^{13}\text{C}^{18}\text{O}$

P. CACCIANI

Laboratoire Aimé Cotton, Campus d'Orsay, 91405 Orsay Cedex, France

W. UBACHS AND P. C. HINNEN

Department of Physics and Astronomy, Vrije Universiteit, De Boelelaan 1081, 1081 HV Amsterdam, The Netherlands

AND

C. LYNGÅ, A. L'HUILLIER, AND C.-G. WAHLSTRÖM

Department of Physics, Lund Institute of Technology, P.O. Box 118, S-22100 Lund, Sweden

Received 1998 February 17; accepted 1998 March 31; published 1998 May 6

ABSTRACT

In a laboratory investigation using tunable picosecond lasers in a pump-probe configuration, lifetimes of the $E\ ^1\Pi$, $v = 0$ and $v = 1$ states of CO are experimentally determined. For $E\ ^1\Pi$, $v = 1$, the lifetime is found to vary significantly for the isotopomers. Combining the data with existing oscillator strengths, predissociation probabilities within the range of 93.6%–98.6% result for $^{12}\text{C}^{16}\text{O}$, $^{13}\text{C}^{16}\text{O}$, and $^{13}\text{C}^{18}\text{O}$ in the case of the $E\ ^1\Pi$, $v = 1$ state, which is the state that predominantly influences the isotopic fractionation of CO in interstellar clouds.

Subject headings: ISM: molecules — molecular data — techniques: spectroscopic

1. INTRODUCTION

Carbon monoxide, the second most abundant molecule in the galaxy and the prototypical molecule for radio-astronomical studies, is a key ingredient in the chemistry of the interstellar medium. In the last decade, the photochemistry of CO and its isotopomers in diffuse and translucent clouds, as well as in photodissociation regions, has been clarified, to some extent, through modeling on the one hand (van Dishoeck & Black 1988; Viala et al. 1988) and laboratory experiments on the other. A number of spectroscopic overview studies (Letzelter et al. 1987; Eidelsberg & Rostas 1990; Stark et al. 1991) in the wavelength range 91.2–110 nm, of interest for the photodissociation of CO in the interstellar medium, has yielded the general result that *predissociation* rather than continuum absorption is the main destruction mechanism for interstellar CO. The fact that predissociation, i.e., dissociation through bound states that are coupled to continuum states, is by its nature strongly wavelength dependent makes the photodissociation of CO a complicated problem. The richness of mutually interacting Rydberg and valence states is an additional complicating factor, particularly because the couplings appear to be strongly isotope dependent (Rostas et al. 1994). Following the first overview studies by classical and synchrotron techniques, narrow-band lasers are being used to unravel spectral features and predissociation phenomena in finer detail (Levelt, Ubachs, & Hogervorst 1992; Eikema, Hogervorst, & Ubachs 1994; Drabbel et al. 1993; Komatsu et al. 1995; Ubachs et al. 1997). The progress via laser-based measurements has led to revised listings of spectral constants and predissociation rates (Ubachs et al. 1994; Morton & Noreau 1994).

In the advanced model of van Dishoeck & Black (1988), the effects of line broadening associated with predissociation, self-shielding of CO lines, shielding by atomic and molecular hydrogen lines, and isotope-selective shielding are examined extensively as a function of the depth in interstellar clouds. These photoinduced phenomena, as well as ion-molecule exchange reactions at low temperatures, result in strong isotope-dependent effects on the photodissociation rates. Thus, in the interior of interstellar clouds, larger effective photodissociation rates are found for the less abundant, heavier isotopomers. In particular, the excitation of the $E\ ^1\Pi$ state at 107.61 nm for

$v = 0$ and 105.17 nm for $v = 1$ seems to be of paramount importance, since these transitions have large oscillator strengths that coincide with a relatively high intensity of the interstellar radiation field. The predissociation probability of the $E\ ^1\Pi$, $v = 1$ state is identified as the decisive parameter governing isotope selectivity in predissociation. In their conclusion, van Dishoeck & Black (1988) propose an independent measurement of the predissociation probabilities of the $E-X$ (0, 0) and (1, 0) bands to further validate their model.

Here we report on direct time-domain measurements of the lifetimes of the $E\ ^1\Pi$, $v = 0$ and $v = 1$ states for the $^{12}\text{C}^{16}\text{O}$, $^{13}\text{C}^{16}\text{O}$, and $^{13}\text{C}^{18}\text{O}$ isotopomers. The harmonic radiation generated by the picosecond laser of the Lund Laser Centre, previously used for lifetime measurements of He (Larsson et al. 1995), is used in a pump-probe configuration to detect time-delayed photoionization. The vacuum ultraviolet (VUV) radiation is tuned to resonance with the $E-X$ (0, 0) or (1, 0) band, while the second laser, at 355 nm and at a variable delay, induces photoionization. Resulting CO^+ ions are accelerated in a DC-electric field, mass-selected in a time-of-flight drift tube, and detected by an electron multiplier. The signal is registered on and read from a digital storage oscilloscope. The data on $^{12}\text{C}^{16}\text{O}$ were obtained from CO gas in natural abundance. The $^{13}\text{C}^{16}\text{O}$ and $^{13}\text{C}^{18}\text{O}$ data are obtained from a 99% ^{13}C -enriched sample of CO in which an amount of approximately 20% $^{13}\text{C}^{18}\text{O}$ is found. Since the bands for the isotopomers overlap, the isotope selectivity of the time of flight is required for distinction between $^{13}\text{C}^{16}\text{O}$ and $^{13}\text{C}^{18}\text{O}$ signals.

2. EXPERIMENTAL METHOD AND RESULTS

A distributed-feedback dye laser (DFDL), tunable in the near-infrared, is pumped by the 532 nm output of an amplified mode-locked Nd-YAG laser, running at a 10 Hz repetition rate. The low-energy pulses of the DFDL are amplified in a titanium-sapphire crystal, which is pumped by a Q -switched Nd-YAG laser. The duration of the resulting infrared pulses is 50–70 ps, and via generation of the seventh harmonic in a pulsed jet of Xe atoms, a VUV pulse is produced with an estimated duration of 30 ps. The UV probe pulse at 355 nm is the third harmonic output of the same Nd-YAG laser, used to pump the dye laser. Its pulse duration was measured with a streak camera to be 60

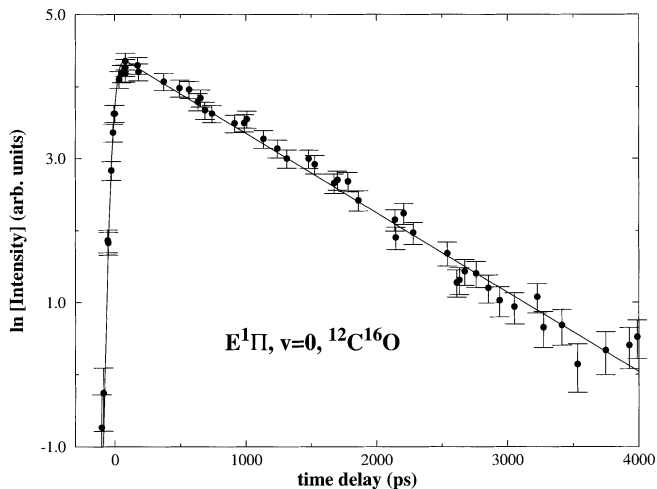


FIG. 1.—Measurement of pump-probe decay transient with the VUV laser set at the Q -branch of the $E^1\Pi, v=0$ state of $^{12}\text{C}^{16}\text{O}$. Note that the vertical axis is on a log scale.

ps. We estimate the instrument-limited temporal resolution to be 80 ps. The optical delay between the VUV and UV pulses is manually set and measured on an optical rail allowing delays of up to 5.7 ns. The VUV beam is focused and crossed at 45° with the loosely focused UV beam. This geometry strongly reduces radiative trapping, an effect that often hinders reliable lifetime measurements. The CO gas is pulsed into the interaction zone with a pulsed valve at 7 cm distance, ensuring that the free-jet expansion reaches the collision-free regime at the interaction zone. The bandwidth of the DFDL laser is 3.5 cm^{-1} in the infrared, resulting in a resolution of 25 cm^{-1} in the VUV, which is insufficient to resolve individual rotational lines. However, a distinction between P -, Q -, and R -branches could be made, while an effect of rotational cooling could be observed from the spectral signature also.

In Figure 1, the data of a pump-probe delay measurement for the $E^1\Pi, v=0$ state of the $^{12}\text{C}^{16}\text{O}$ isotopomer are presented. Each data point is an average over the signal obtained from 200 laser pulses and is plotted on a logarithmic scale. The rising edge at negative delays of the probe laser is the signature of the limited temporal resolution of the instrument. This is accounted for in a fitting routine by convolving an exponential decay function with a Gaussian profile of 80 ps FWHM. The uncertainty in the measured intensity at each delay setting is composed of different contributions, reading uncertainties on the digital oscilloscope, depending on magnification and background noise, as well as shot-to-shot intensity fluctuations of the laser pulses. The latter uncertainties are on the order of 30%–50% for the VUV-picosecond laser. Even with averaging over 200 shots, the intensity fluctuations remain the main source of error at short delays. At longer delays, the background noise becomes significant. In a weighted fit, a constant and an intensity-dependent term are incorporated in an expression for the error estimate for each data point, where the intensity-dependent noise scales as the square root of the intensity, as is appropriate for photon counting noise. As a consequence, the data at longer delays receive less weight in the fit and appear with larger errors in the log plot of Figure 1. For the decays of the $E^1\Pi, v=0$ state of $^{12}\text{C}^{16}\text{O}$, careful checks were made to investigate possible systematic effects due to the variation of the density in the molecular beam, the UV intensity of the probe laser, and the timing of the laser pulses with respect to

TABLE 1

MEASURED LIFETIMES τ (IN UNITS OF PICOSECONDS) ^a			
$E^1\Pi$	$^{12}\text{C}^{16}\text{O}$	$^{13}\text{C}^{16}\text{O}$	$^{13}\text{C}^{18}\text{O}$
$v=0$	912 ± 60	1350 ± 200	1250 ± 200
$v=1$	118 ± 20	198 ± 25	270 ± 45

^a For the $E^1\Pi, v=0$ and $v=1$ states of CO isotopomers.

the opening of the valve that delivers the molecular jet. From spectral analyses, it follows that the rotational distribution is colder earlier in the gas pulse. No dependence of the obtained lifetime on pressure, UV intensity, or rotational temperature is found, for either the (e) or the (f) parity state in $E^1\Pi, v=0$; these states were independently probed by setting the laser on the Q - or P/R -branch. In Table 1, the resulting values for the lifetimes are listed with an estimated error. Although the error margins on the lifetimes for the ^{13}CO isotopomers are substantial, we find a definite indication of longer lifetimes for the ^{13}C containing isotopomers.

In Figure 2, a pump-probe decay curve for the $E^1\Pi, v=1$ state of $^{13}\text{C}^{16}\text{O}$ is presented with the VUV radiation tuned to the Q -branch. Here the data show a much faster decay than in the case of $v=0$. In view of the shorter lifetime and the weaker transition strength, both collisional effects and possible radiative trapping phenomena are less important. The systematic issue to address for the case of $v=1$ is the deconvolution of the 80 ps instrument width. From a variation of the convolution parameter, it is found that indeed the estimated 80 ps leads to a minimum standard deviation in the fitting routine. The lifetimes obtained are listed in Table 1. In case of the $E^1\Pi, v=1$ state, the lifetime of the main $^{12}\text{C}^{16}\text{O}$ isotopomer is found to be significantly shorter than that of $^{13}\text{C}^{16}\text{O}$ and $^{13}\text{C}^{18}\text{O}$.

3. DISCUSSION

Hitherto, only one lifetime measurement has been reported on the $E^1\Pi$ state, the one on the $v=0$ level by Smith (1978), which yields a value of $\tau_0 = 1.5 \pm 0.2$ ns, slightly higher than the present value. Eidelsberg & Rostas (1990) give some rather crude estimates, based on fluorescence yield measurements. However, their results of 10^{-9} s for $E^1\Pi, v=0$ and 10^{-10} s

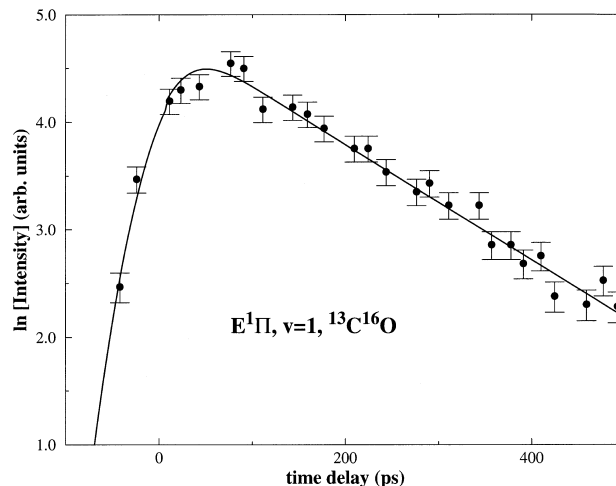


FIG. 2.—Lifetime measurement of the $E^1\Pi, v=1$ state of $^{13}\text{C}^{16}\text{O}$ with the laser set at the Q -branch. The slow rising edge is a consequence of the finite time resolution.

for $E^1\Pi$, $v = 1$ closely match the present dedicated measurements.

An astrophysically important property is the competition between radiative and dissociative decay on excitation. The predissociation probabilities can be derived from the combined results of the present measurements and previously obtained oscillator strengths for the $E-X$ system. Following Morton & Noreau (1994), the oscillator strength $f_{v'v''}$ is related to a decay rate $A_{v'v''}$ via

$$\frac{1}{2}f_{v'v''} = \left(4\pi\epsilon_0 \frac{m_e c}{8\pi^2 e^2}\right) \lambda_{v'v''}^2 A_{v'v''}, \quad (1)$$

where the factor of $\frac{1}{2}$ represents the twofold Λ -doublet degeneracy of the upper state $E^1\Pi$. For clarity, we note that in this case, $f_{v'v''}$ represents the total oscillator strength integrated over the P -, Q -, and R -branches of a $E-X$ (v' , v'') band. The value of the constant in parentheses can be evaluated to $1.4992 \times 10^4 \text{ m}^{-2} \text{ s}$. In the literature, a variety of oscillator strengths have been published. For the $E-X$ (0, 0) band, f_{00} values range from 0.0365 (Eidelsberg & Rostas 1990) to 0.071 obtained in two independent electron-scattering experiments (Chan, Cooper, & Brion 1993; Ciocca, Kanik, & Ajello 1997). An intermediate value of 0.049 by Stark et al. (1992), obtained from high-resolution synchrotron radiation absorption, coincides with a value from ab initio calculations (Kirby & Cooper 1989). Since the purely radiative lifetime τ_0 of the upper level $E^1\Pi$, $v = 0$ corresponds to a summation of partial decay rates,

$$\frac{1}{\tau_0} = A_0 = \sum_{X, v''} A_{0v''} + \sum_{A, B, v''} A_{0v''}, \quad (2)$$

an assumption has to be made on the decay in vibrational levels of the electronic ground state other than $X^1\Sigma^+$, $v'' = 0$. For the absorptive transition originating in $v'' = 0$, it is found that 94% of the oscillator strength is in the (0, 0) band (Eidelsberg & Rostas 1990; Kirby & Cooper 1989; Stark et al. 1992), and by applying sum rules, the same follows for the emission. The radiative decay into the $A^1\Pi$ and $B^1\Sigma^+$ electronic states, amounting to 8% (Kirby & Cooper 1989), should be included. This leads to a total rate A_0 , which is 1.16 times the partial rate A_{00} . The predissociation probability η then follows:

$$\eta_{\text{pre}} = 1 - A_0\tau, \quad (3)$$

where τ is the true lifetime of the upper state, as measured in the present study. Using a number of published f_{00} values, and deriving A_0 from these, the predissociation probabilities can be calculated. The results are listed in Table 2. For the $E^1\Pi$, $v = 1$ level, the same value for A_0 can be used, i.e., $A_1 = A_0$, since the total radiative decay of a vibrational series is constant to a very good approximation.

The value for the predissociation probability of the $E^1\Pi$, $v = 0$ state is found to vary in the range 69%–89%. Since the uncertainty in the lifetime measurements is a minor part of the uncertainty in η , we have not quoted errors in Table 2. Improved values for predissociation probabilities await the determination

TABLE 2
PREDISSOCIATION PROBABILITIES η^a

$E^1\Pi$	CO Isotopomer	η_s (%)	η_E (%)	η_C (%)
$v = 0$	$^{12}\text{C}^{16}\text{O}$	85.1	88.9	78.4
	$^{13}\text{C}^{16}\text{O}/^{13}\text{C}^{18}\text{O}$	78.7	84.2	69.2
$v = 1$	$^{12}\text{C}^{16}\text{O}$	98.1	98.6	97.2
	$^{13}\text{C}^{16}\text{O}$	96.8	97.6	95.3
	$^{13}\text{C}^{18}\text{O}$	95.6	96.7	93.6

NOTE.—The values η_s and η_E pertain to predissociation probabilities that were obtained by using the oscillator strengths of Stark et al. 1992 and Eidelsberg & Rostas 1990, respectively. The value η_C is obtained when using the oscillator strength 0.071 (Chan et al. 1993; Ciocca et al. 1997). In view of the small discrepancies in lifetimes between $^{13}\text{C}^{16}\text{O}$ and $^{13}\text{C}^{18}\text{O}$ for the $v = 0$ levels, the average predissociation probability for these isotopomers is listed. The uncertainties in the predissociation probabilities can be determined from the uncertainties in the measured f -values.

^a For the $E^1\Pi$, $v = 0$ and $v = 1$ states of CO isotopomers.

of more accurate, i.e., more reliable, oscillator strengths. In a recent study by Ciocca et al. (1997), direct electron-impact-induced emission from the $E^1\Pi$, $v = 0$ state was observed at high resolution. A higher fluorescence cross section for the Q -branch indicates a lower predissociation probability on the $E^1\Pi$ -(f) parity components. In the present study, this parity dependence is not found. For this purpose, a lifetime measurement was performed with the laser set on the P -branch, at a wavelength position where, even in view of the large bandwidth of the VUV laser, no signal is obtained from the blue-degraded Q -branch in the (0, 0) band.

An interesting conclusion from a molecular physics point of view is that the lifetime of the $E^1\Pi$, $v = 1$ state is strongly dependent on the specific isotopomer. The reason for this effect is not yet clear. In two studies (Baker et al. 1993; Cacciani, Hogervorst, & Ubachs 1995), it was determined that a $k^3\Pi$ state causes an accidental predissociation near $J = 9$ for $^{12}\text{C}^{16}\text{O}$ and at a different J -value for the other isotopomers. Most likely, there is also an additional effect due to a coupling with a repulsive continuum that causes the predissociation. From an astrophysics viewpoint, these isotope-dependent effects are not decisive since it is firmly established in the present study that for the $E^1\Pi$, $v = 1$ state, for all isotopomers and for all oscillator strengths existing in the literature, the predissociation probability remains in the range 93.6%–98.6%, even when the large spread in published oscillator strengths is considered. In fact, the major assumption of van Dishoeck & Black (1988) that the predissociation does not significantly deviate from 100% is validated in the present experimental study.

P. C., P. H., and W. U. wish to thank the Lund Laser Centre for their hospitality. This work was supported by the European Community via the TMR-grant “Access to Large Scale Facilities,” contract ERBFMGECT950020(DG12). We acknowledge the support from the Swedish Natural Science Research Council. We thank A. Persson for technical support, S. Svanberg for stimulating discussions, and D. Morton (Herzberg Institute) for clarification of equation (1) and for critically reading the manuscript.

REFERENCES

- Baker, J., Lemaire, J. L., Couris, S., Vient, A., Malmasson, D., & Rostas, F. 1993, *Chem. Phys.*, 178, 569
Cacciani, P. C., Hogervorst, W., & Ubachs, W. 1995, *J. Chem. Phys.*, 102, 8308

- Chan, W. F., Cooper, G., & Brion, C. E. 1993, *Chem. Phys.*, 170, 123
Ciocca, M., Kanik, I., & Ajello, J. 1997, *Phys. Rev. A*, 55, 3547
Drabbels, M., Heinze, J., ter Meulen, J. J., & Meerts, W. L. 1993, *J. Chem. Phys.*, 99, 5701

- Eidelsberg, M., & Rostas, F. 1990, *A&A*, 235, 472
- Eikema, K. S. E., Hogervorst, W., & Ubachs, W. 1994, *Chem. Phys.*, 181, 217
- Kirby, K., & Cooper, D. L. 1989, *J. Chem. Phys.*, 90, 4895
- Komatsu, M., Ebata, T., Maeyama, T., & Mikami, N. 1995, *J. Chem. Phys.*, 103, 2420
- Larsson, J., Mevel, E., Zerne, R., L'Huillier, A., Wahlström, C.-G., & Svanberg, S. 1995, *J. Phys. B*, 28, L53
- Letzelter, C., Eidelsberg, M., Rostas, F., Breton, J., & Thieblemont, B. 1987, *Chem. Phys.*, 114, 273
- Levelt, P. F., Ubachs, W., & Hogervorst, W. 1992, *J. Chem. Phys.*, 97, 7160
- Morton, D. C., & Noreau, L. 1994, *ApJS*, 95, 301
- Rostas, F., Launay, F., Eidelsberg, M., Benharrou, M., Blaess, C., & Huber, K. P. 1994, *Canadian J. Phys.*, 72, 913
- Smith, W. H. 1978, *Phys. Scr.*, 17, 513
- Stark, G., Smith, P. L., Ito, K., & Yoshino, K. 1992, *ApJ*, 395, 705
- Stark, G., Yoshino, K., Smith, P. L., Ito, K., & Parkinson, W. H. 1991, *ApJ*, 369, 574
- Ubachs, W., Eikema, K. S. E., Hogervorst, W., & Cacciani, P. C. 1997, *J. Opt. Soc. Am. B*, 14, 2469
- Ubachs, W., Eikema, K. S. E., Levelt, P. F., Hogervorst, W., Drabbels, M., Meerts, W. L., & ter Meulen, J. J. 1994, *ApJ*, 427, L55
- van Dishoeck, E. F., & Black, J. H. 1988, *ApJ*, 334, 771
- Viala, Y. P., Letzelter, C., Eidelsberg, M., & Rostas, F. 1988, *A&A*, 193, 265