# A laser system for the spectroscopy of highly charged bismuth ions 

S. Albrecht • S. Altenburg • C. Siegel • N. Herschbach • G. Birkl

Published online: 10 September 2011
© Springer-Verlag 2011


#### Abstract

We present and characterize a laser system for the spectroscopy on highly charged ${ }^{209} \mathrm{Bi}^{82+}$ ions at a wavelength of 243.87 nm . For absolute frequency stabilization, the laser system is locked to a near-infra-red laser stabilized to a rubidium transition line using a transfer cavity based locking scheme. Tuning of the output frequency with high precision is achieved via a tunable rf offset lock. A sample-and-hold technique gives an extended tuning range of several THz in the UV. This scheme is universally applicable to the stabilization of laser systems at wavelengths not directly accessible to atomic or molecular resonances. We determine the frequency accuracy of the laser system using Dopplerfree absorption spectroscopy of $\mathrm{Te}_{2}$ vapor at 488 nm . Scaled to the target wavelength of 244 nm , we achieve a frequency uncertainty of $\sigma_{244 \mathrm{~nm}}=6.14 \mathrm{MHz}$ (one standard deviation) over six days of operation.


## 1 Introduction

The spectroscopic investigation of hydrogen-like atomic systems has developed into an important test bed for research on atomic structure and bound-state quantum electrodynamics (QED). Already high-precision measurements on atomic hydrogen confirm QED theory and its predictions to a high degree. Even more accurate tests of the predictive power of QED can be achieved by increasing either the precision of the measurement or by increasing the contribution of the effect to be measured [1]. The use of highly

[^0]charged heavy ions shifts the magnitude of QED effects from the perturbative regime to a region where higher order terms become important and perturbation theory is no longer valid [2]. In addition, the energy scale of the $1 s$ ground state hyperfine structure is shifted to the laser accessible regime above a nuclear charge number of $Z \approx 60$ and the upper state lifetime enters a regime where acceptable fluorescence rates are expected [3].

Based on these considerations, the spectroscopic investigation of the M1 transition of the ground state hyperfine structure of hydrogen-like bismuth $\left({ }^{209} \mathrm{Bi}^{82+}\right)$ at 243.87 nm has become one of the prime targets for further research along these lines. Theoretical investigations of the hyperfine transition energy, the lifetime and their respective QED contributions have been performed [4-6]. Measurements of the transition energy have been realized on bunches of ${ }^{209} \mathrm{Bi}^{82+}$-ions traveling at a velocity of 0.6 c in the experimental storage ring ESR at the Gesellschaft für Schwerionenforschung (GSI) [4, 7]. The laboratory value of the transition wavelength was determined to 243.87 (4) nm [7]. The uncertainty was dominated by the limitation in the velocity calibration of the fast moving ions. This initial uncertainty of approximately 200 GHz has been reduced to approximately 100 GHz in a later analysis [4]. To overcome this limitation, the deceleration of highly charged ions and a spectroscopic investigation of trapped ions have been proposed [3, 8]. Spectroscopy of highly charged ions stored in Penning traps is expected to reach accuracies which make tests of bound-state QED calculations significantly more stringent and give access to improved measurement of fundamental quantities [9].

At the SPECTRAP experiment at the HITRAP facility [8] at GSI, systematic measurements on several species of highly charged hydrogen- and lithium-like ions will improve the spectroscopic resolution by up to three orders of magni-
tude over previous experiments. The assembly consists of a cryogenic Penning trap, a frequency stabilized laser for excitation of the hyperfine transition under investigation, and a detection system for emitted fluorescence light [3]. The trap will be loaded with highly charged ions produced at the GSI accelerator facility which are decelerated and delivered by the HITRAP beam-lines and cooled to liquid helium temperatures inside the Penning trap. For efficient spectroscopy of ${ }^{209} \mathrm{Bi}^{82+}$ the laser system should produce laser light with a power of several mW at 243.87 nm and a tuning range significantly larger than the standard deviation of 100 GHz of the previous measurement. In addition, the laser frequency should be stable to a precision and accuracy comparable to or below the expected Doppler-width of the transition of the ions inside the Penning trap which is expected to be approximately 30 MHz .

## 2 Laser design and stabilization scheme

We have developed, built, and characterized a laser systems which fulfills these requirements. The system consists of four major components (Fig. 1): a commercial frequencyquadrupled diode laser system (target laser, Toptica TAFHG 110) generating light at a wavelength of 243.87 nm , a pair of offset-locked diode lasers operating at 780 nm (master and transfer laser), a confocal transfer cavity locking the target laser to the transfer laser at a large wavelength offset, and a set of spectroscopic diagnostics at 780 nm (rubidium $(\mathrm{Rb})$ absorption cell) and 488 nm (tellurium $\left({ }^{130} \mathrm{Te}_{2}\right)$


Fig. 1 Schematic overview of the laser system. The master laser, stabilized to a rubidium transition, serves as absolute frequency reference. The transfer laser is offset-locked to the master laser by an rf beat measurement. The transfer cavity is stabilized to the transfer laser and serves as reference for locking the target laser. Frequency doubling and quadrupling generates light at 488 and 244 nm . In the inset (gray frame) the transmission signal (top) and the calculated dispersive signal (bottom) of the transfer cavity are shown indicating the capture range $(R)$ of about 850 MHz
absorption cell) for characterization of the different components of the frequency stabilization chain.

The target laser produces light around 244 nm and an output power of up to 15 mW . This is achieved by fourth harmonic generation of the light of a tapered amplifier with an output power of 800 mW seeded by an external cavity diode laser operating at a wavelength of 976 nm . The manufacturer specifies the linewidth of the system to be smaller than 4 MHz at 244 nm ( $1-\mu \mathrm{s}$ measurement interval). The modular setup with two successive frequency doubling stages gives access to three laser fields: light at 976 nm is used for frequency stabilization, light at 488 nm is used for frequency diagnostics on ${ }^{130} \mathrm{Te}_{2}$ vapor, and light at 244 nm is used for high-resolution spectroscopy of ${ }^{209} \mathrm{Bi}^{82+}$. In order to be able to implement the required large tuning range of several 100 GHz , the direct stabilization of one of these wavelengths to an atomic or molecular transition is not possible. For that reason, we implemented a stabilization scheme where the fundamental laser output at 976 nm is locked to a lengthstabilized transfer cavity. The transfer cavity serves as a stable reference similar to the one used in [10], where a cavity is actively stabilized to a polarization stabilized helium-neon laser. In contrast to this approach, our transfer cavity is stabilized to a tunable diode laser (transfer laser). By tuning the frequency of the transfer laser, we can tune the resonance frequency of the transfer cavity and in this manner also the output frequency of the target laser. An overview of the involved laser wavelengths is provided by Fig. 2: The anchor of the stabilization chain is the master laser stabilized to a ${ }^{85} \mathrm{Rb}$ transition at 780 nm by Doppler-free saturation spectroscopy. The transfer laser is offset-locked to the master laser with a computer-controlled frequency offset in the range of 2 to 10.5 GHz with respect to 780 nm . The transfer cavity is stabilized to the transfer laser, bridges the wavelength gap to 976 nm , and serves for locking of the fundamental output of the target laser system. Using a sample-and-hold technique (see Sect. 2.3) a scan range of


Fig. 2 Overview in the output frequencies occurring in the frequency stabilized laser system for spectroscopy of hydrogen-like bismuth ${ }^{209} \mathrm{Bi}^{82+}$
several 100 GHz can be achieved. Finally, two successive frequency doubling stages generate light at 488 and 244 nm .

The following sections give a more detailed discussion of the stabilization chain.

### 2.1 Rb-stabilization and RF-offset-lock

The master and transfer lasers are home-built externalcavity diode lasers operating at a wavelength of 780 nm . The master laser is stabilized to the $5^{2} S_{1 / 2}(F=3) \rightarrow$ $5^{2} P_{3 / 2}\left(F^{\prime}=3, F^{\prime}=4\right)$ cross-over resonance of ${ }^{85} \mathrm{Rb}$ by Doppler-free absorption spectroscopy in a mu-metal shielded vapor cell. The stabilization is based on phasesensitive detection of the saturated absorption signal using a lock-in amplifier with direct modulation of the diode laser's injection current at a frequency of 70 kHz . The feedback bandwidth of 10 kHz of the PI servo loop is defined by the time constant of the lock-in amplifier. Typical short-term line-widths of the 780 nm lasers are in the range of 50 kHz .

A heterodyne beat-signal between the outputs of master and transfer laser is mixed with the output of an rfsynthesizer with a double balanced mixer. A frequency-tovoltage converter produces an error signal proportional to the frequency offset between the beat and the synthesizer frequency. This error signal is fed to a PI controller for stabilization of the transfer laser frequency. The transfer laser frequency can be tuned with respect to the fixed master laser frequency with sub-kHz resolution by tuning the synthesizer output. This allows us to scan the frequency of the transfer laser with high accuracy and in an arbitrary frequency sequence.

The transfer laser stabilization and tunability are enhanced using the intra-cavity stabilization scheme described in [11]. Here, the internal cavity of the laser diode consisting of its front and rear surface is mode-matched to an external cavity formed by the rear surface of the laser diode and an external cavity mirror. Mode-matching is controlled by the injection current of the laser diode. With this technique we reach a mode-hop free tuning range of 8.5 GHz (at 780 nm ), limited by the bandwidth of the beat-note detection.

We use an independent Doppler-free absorption spectroscopy on rubidium to characterize the performance of the frequency stabilization and to determine the uncertainty of the transfer laser frequency. By repeated scans over the Doppler-free resonances of the $5^{2} S_{1 / 2}(F=1) \rightarrow$ $5^{2} P_{3 / 2}\left(F^{\prime}\right)$ transitions of ${ }^{87} \mathrm{Rb}$ and by fitting Lorentzian profiles to the occurring resonance lines (Fig. 3(a)) we can extract the center frequencies of the observed lines. The resulting histograms of the center frequencies of all lines and of the $(F=1) \rightarrow\left(F^{\prime}=1, F^{\prime}=2\right)$ cross-over transition are presented in Figs. 3(b) and 3(c), respectively. A Gaussian fit (dashed curve in Fig. 3(c)) gives a statistical uncertainty


Fig. 3 (a) Doppler-free absorption spectroscopy on ${ }^{87} \mathrm{Rb}$ with fitted Lorentzian profiles; (b) Histogram of the extracted center frequencies; (c) Histogram of the center frequencies of the ${ }^{87} \mathrm{Rb}$ $(F=1) \rightarrow\left(F^{\prime}=1, F^{\prime}=2\right)$ cross-over transition. The standard deviation ( $\sigma_{780 \mathrm{~nm}}=83 \mathrm{kHz}$ ) of the fitted Gaussian function gives a value for the frequency uncertainty
of $\sigma_{780 \mathrm{~nm}}=83 \mathrm{kHz}$ (one standard deviation) for data accumulated over several hours and $\sigma_{780 \mathrm{~nm}}=123 \mathrm{kHz}$ for data compiled over several days.

### 2.2 Transfer cavity

We use a transfer cavity for stabilization of the target laser to the transfer laser with large wavelength difference. The length of the transfer cavity is locked to the transfer laser wavelength using a piezo transducer in one of the cavity mirror mounts. Our locking scheme, which includes a sample-and-hold circuit for a controlled change of the cavity mode number (see Sect. 2.3) is applicable to a wide range of target wavelengths and in addition allows us to extend the tuning range of the target laser well beyond the transfer laser tuning range. The transfer cavity consists of a temperature-stabilized aluminum spacer and low-reflective silver mirrors giving a free spectral range (FSR) of about 2 GHz . An aluminum spacer is sufficient for our purpose due to the active length stabilization to the transfer laser. In a later stage switching to Invar, Zerodur, or ULE may be considered. The free spectral range is measured by scanning the laser frequency between absolutely calibrated features of tellurium (Sect. 3.1) and recording the number of transmission fringes. These measurements resulted in a $F S R=$ 2002.345 (13) MHz. By changing the cavity length to the next mode number, the $F S R$ is changed by approximately 40 kHz .

The reflectivity of the cavity mirrors with coatings of 15 nm Ag and 40 nm MgF 2 was chosen to be $69 \%$ at 976 nm
and $70 \%$ at 780 nm , respectively, in order to give a large capture range for the transfer cavity locks. The theoretical value for the finesse of 8.5 is confirmed experimentally by a measured value of $8.24(F W H M=243 \mathrm{MHz})$. Though the large fringe width makes the system sensitive to offset drifts in the electronics, monitoring and correcting these drifts resulted in insignificant contributions to the frequency drifts. The temperature stabilization of the cavity spacer has small residual thermal variations resulting in piezo transducer stabilization voltage variations on long time-scales. We determined that these variations of approximately 30 V translate to a remaining temperature fluctuation of about $0.2^{\circ} \mathrm{C}$ which corresponds to the expected precision of the temperature stabilization used.

The length stabilization of the transfer cavity as well as the stabilization of the target laser to the transfer cavity are done by top-of-fringe locking since it is insensitive to intensity fluctuations. By modulating the position of one of the cavity mirrors at 5 kHz we produce the error signals for both stabilization circuits using phase-sensitive demodulation with lock-in amplifiers.

### 2.3 Stabilization of target laser system

We stabilize the fundamental wavelength of the target laser system at 976 nm to the transfer cavity. A sample-and-hold technique allows us to extend the tuning range of the target laser beyond the tuning range of the transfer laser: at the limit of the tuning range, the feedback controller is muted by external control. This causes the control voltage to stay constant ("hold operation") on the last value before muting and the target laser to hold its set-point. The wavelength is then held without feedback, but small wavelength drifts may occur. Next we scan the transfer cavity back by an integer multiple of the $F S R$ and relock the target laser to another cavity fringe. The large capture range due to the small finesse of the cavity (see inset in Fig. 1) guarantees relocking although the target laser wavelength may have a small deviation from the expected wavelength. The fringe shifting by sample-and-hold ensures the cavity length on average to stay the same and the FSR to stay on a well defined value. In addition, the mode-hop free tuning range of the target laser system of approximately 40 GHz at 244 nm (limited by the finite tuning range of the doubling and quadrupling cavities) can be fully exploited. Realigning the doubling and quadrupling stages results in a tuning range of the system of more than 100 GHz in the UV as required.

## 3 Results and discussion

### 3.1 Spectroscopic measurements on tellurium

The fundamental frequency of the target laser system is frequency doubled to 488 nm and then again frequency dou-
bled to 244 nm by two separate external doubling cavities. A small fraction of the blue laser light at 488 nm is diverted for Doppler-free saturation spectroscopy in a vapor cell of tellurium $\left({ }^{130} \mathrm{Te}_{2}\right)$. Similar to iodine, ${ }^{130} \mathrm{Te}_{2}$ has many resonance lines that cover most of the visible spectrum, so that the second harmonic frequency of the target laser can be used to determine the frequency uncertainty of the stabilized laser system. In addition, high-precision reference lines in ${ }^{130} \mathrm{Te}_{2}$ can be used to give absolute accuracy to our laser system. A $10 \mathrm{~cm}{ }^{130} \mathrm{Te}_{2}$ cell similar to the one employed in [12-14], is heated to $500^{\circ} \mathrm{C}$ in a tube furnace. By scanning the laser system, we record the Doppler-broadened and the Doppler-free transition lines of ${ }^{130} \mathrm{Te}_{2}$ over a range of several 100 GHz . We can identify numerous Doppler-broadened lines of the "tellurium atlas" [15] and the Doppler-free lines published in [12-14, 16-18]. In specific, we scan the known $\mathrm{d}_{4}$ reference line [12] numerous times (Fig. 4(a)) and extract the center frequency of this line, by fitting a Lorentz function to each measurement. The extracted center frequencies are plotted in a histogram (Fig. 4(b)). A Gaussian fit function describes the distribution well. The standard deviation of the Gaussian is $\sigma_{488} \mathrm{~nm}=3.07 \mathrm{MHz}$ (at 488 nm ) for the compiled data from over 1300 measurements recorded in six consecutive days including a 50 hour period of continuous laser operation and data taking. For the final target wavelength of 244 nm this corresponds to an uncertainty of $\sigma_{244} \mathrm{~nm}=6.14 \mathrm{MHz}$ which is well below the required uncertainty of 30 MHz as given by the Doppler broadening of the targeted ${ }^{209} \mathrm{Bi}^{82+}$ transition.

### 3.2 Influence of refractive index of air

During the six day measurement run resulting in the data of Fig. 4(b), we observed a drift in the measured center frequency of a few MHz , which we attribute to a change in air pressure. Using this additional information it is possible to further reduce the uncertainty of the target laser output frequency in our open cavity configuration. Ultimately, housing the transfer cavity in a constant-pressure or evacuated environment should eliminate this effect.

To gain further insight, we analyzed our data in the light of potential temperature and pressure dependences of the target laser wavelength caused by a wavelength dependent variation of the transfer cavity $F S R$ at 780 nm and 976 nm . A good description of the influence of pressure and temperature on an open transfer cavity is provided by [10]. The resonance frequency of the transfer cavity for the $T E M_{00}$ mode is given by
$v_{N}=\frac{N c}{2 n d}$
where $\nu_{N}$ is the resonance frequency, $N$ is the order of the longitudinal mode, $c$ is the vacuum speed of light, $d$ is the


Fig. 4 (a) Doppler-broadened (blue) and Doppler-free (red) spectrum of tellurium around the $\mathrm{d}_{4}$ reference feature. The dashed line indicates the fitted Lorentzian profile of the $\mathrm{d}_{4}$ line. (b) Histogram of the center frequencies extracted from repeated measurements of the $d_{4}$ line. The data were taken over six days resulting in 1300 individual measurements. The dashed line represents a Gaussian fit with a standard deviation of $\sigma_{488} \mathrm{~nm}=3.07 \mathrm{MHz}$, giving $\sigma_{244} \mathrm{~nm}=6.14 \mathrm{MHz}$
separation of the cavity mirrors and $n$ is the index of refraction of the medium between the mirrors. At the transfer laser wavelength, changes in the refractive index are compensated by keeping the optical path length between the mirrors constant through the feedback loop. The stabilized frequency of the target laser, on the other hand is affected by the dispersion of air, especially when large frequency spans are bridged by the transfer cavity. The refractive index of air under standard conditions $\tilde{n}$ is calculated for 780 nm and 976 nm following the procedure given in [19] relative to the refractive index of vacuum which is $\tilde{n}_{\text {vac }} \equiv 1$ :
$\tilde{n}(\lambda)-1=\left[8342.13+\frac{2406030}{130-\lambda^{-2}}+\frac{15997}{38.9-\lambda^{-2}}\right] \times 10^{-8}$
with the wavelength $\lambda$ (in $\mu \mathrm{m}$ ). The difference in the refractive indices results to $\tilde{n}_{780}-\tilde{n}_{976}=\delta \tilde{n}=9.3065 \times$ $10^{-7}$. The dependence of the refractive index of air $n$ on pressure $p$ (in Pa ) and temperature $T\left(\right.$ in ${ }^{\circ} \mathrm{C}$ ) is given by
$n(p, T, \tilde{n}(\lambda))-1=(\tilde{n}(\lambda)-1) \cdot \frac{1.0413 \times 10^{-5} \cdot p}{1+3.671 \times 10^{-3} \cdot T}$
according to [19] and can be approximated by a Taylor expansion up to linear order in $T$ around $T_{0}=20^{\circ} \mathrm{C}$
through

$$
\begin{align*}
n(p, T, \tilde{n}(\lambda))-1 \approx & (\tilde{n}(\lambda)-1) \cdot\left(1.0413 \times 10^{-5} \cdot p\right) \\
& \times\left(0.93160-3.1860 \times 10^{-3} \cdot \Delta T\right) \tag{4}
\end{align*}
$$

with $\Delta T=T-T_{0}$. According to (1), the resulting deviation of the target laser frequency is given by:
$\Delta v=v_{N} \cdot \frac{\Delta n}{n}$
For a calculation of the uncertainty in $n$, the linearized Taylor expansion (4) is applied to realistic parameters ( $\Delta p \leq$ $10 \mathrm{hPa} ; \Delta T \leq 0.2^{\circ} \mathrm{C}$ ). Only the difference in the index of refraction $\tilde{n}_{780}-\tilde{n}_{976}=\delta \tilde{n}$ which is not compensated by the cavity length stabilization has to be taken into account resulting in:
$\Delta n(p, T, \delta \tilde{n})=\sqrt{\left(\frac{\partial n}{\partial p} \Delta p\right)^{2}+\left(\frac{\partial n}{\delta T} \Delta T\right)^{2}}$,
with

$$
\begin{align*}
\frac{\partial n}{\partial p} \Delta p & =\delta \tilde{n} \cdot 1.0413 \times 10^{-5} \Delta p \cdot 0.93160 \\
& =9.0280 \times 10^{-12} \Delta p  \tag{7}\\
\frac{\partial n}{\partial T} \Delta T & =\delta \tilde{n} \cdot 1.0413 \times 10^{-5} \cdot p \cdot\left(-3.1860 \times 10^{-3} \Delta T\right) \\
& =-3.0875 \cdot 10^{-9} \Delta T \tag{8}
\end{align*}
$$

using $p=1000 \mathrm{hPa}$. For the assumed temperature and pressure fluctuations of $\Delta T \leq 0.2^{\circ} \mathrm{C}$ and $\Delta p \leq 10 \mathrm{hPa}$, respectively, the contribution of temperature fluctuations can be neglected. Pressure fluctuations result in a change of the stabilized target laser frequency of 0.27 MHz per hPa at 976 nm which scales to 0.54 MHz and 1.08 MHz per hPa at 488 nm and 244 nm , respectively. For the largest assumed pressure variations of 10 hPa , the resulting change in the stabilized target frequency of 10.8 MHz at 244 nm is still well below the targeted absolute accuracy of 30 MHz .

We use our data taken over a six day period with significantly varying air pressure to confirm this dependence and to show that knowledge of the air pressure can be used to further reduce the frequency uncertainty of our laser system. Figure 5 presents the frequency to resonantly excite the $\mathrm{d}_{4}$-line averaged for each day of data taking as a function of the average air pressure reported for the Darmstadt area. We see a clear linear dependence of the $\mathrm{d}_{4}$ frequency on air pressure. Fitting a linear function to the data gives a measured slope of $\frac{\Delta v}{\Delta p}=-0.58(0.11) \mathrm{MHz} / \mathrm{hPa}$ at 488 nm . This slope agrees well with the calculated slope of $0.54 \mathrm{MHz} / \mathrm{hPa}$


Fig. 5 Measured frequency to resonantly excite the tellurium $d_{4}$-line averaged for each day of data taking as a function of the average air pressure for each day. The linear dependence matches the theoretical prediction based on the pressure dependence of the difference in the refractive indices between 780 nm and 976 nm
when taking into account that an increase in the target laser frequency for increasing air pressure has to be compensated by a decrease in the transfer laser frequency to be still in resonance with the selected tellurium line.

## 4 Conclusion

In this article, we have presented a laser system for the high-precision measurement of the hyperfine structure of hydrogen-like ${ }^{209} \mathrm{Bi}^{82+}$. We have discussed a generally applicable approach for stabilization of a laser system to almost any target wavelength by making use of a rubidiumstabilized laser and a broadband transfer cavity. The performance that can be reached depends on the frequency difference between the laser used to stabilize the transfer cavity and the laser to be stabilized to the cavity due to the pressure dependence of the refractive index of air. This influence can be further reduced by monitoring the air pressure or by placing the transfer cavity inside a constant-pressure or evacuated environment. Doppler-free saturation spectroscopy on tellurium can be used to determine the frequency uncertainty of the laser system and to provide reference lines for an absolute frequency calibration of the laser system.

Acknowledgements We thank M. Schlosser, T. Lauber, J. Küber, J. Schütz, S. Tichelmann and the SPECTRAP collaboration for numerous helpful discussions. This work was supported in part by the BMBF (contract numbers 06DA9019I and 06DA9020I).

## References

1. T. Stöhlker, A. Gumberidze, M. Trassinelli, V. Adrianov, H.F. Beyer, S. Kraft-Bermuth, A. Beile, P. Egelhof, The FOCAL collaboration, Lect. Notes Phys. 745, 157 (2008)
2. D.F.A. Winters, M. Vogel, D.M. Segal, R.C. Thompson, W. Nötershäuser, Can. J. Phys. 85, 403 (2007). arXiv: 0704.0560
3. M. Vogel, D.F.A. Winters, D.M. Segal, R.C. Thompson, Rev. Sci. Instrum. 76, 103102 (2005)
4. S. Borneis, A. Dax, T. Engel, C. Holbrow, G. Huber, T. Kühl, D. Marx, P. Merz, W. Quint, F. Schmitt, P. Seelig, M. Tomaselli, H. Winter, K. Beckert, B. Franzke, F. Nolden, H. Reich, M. Steck, Hyperfine Interact. 127, 305 (2000)
5. V.M. Shabaev, A.N. Artemyev, V.A. Yerokhin, O.M. Zherebtsov, G. Soff, Phys. Rev. Lett. 86, 3959 (2001)
6. M. Finkenbeiner, B. Fricke, T. Kühl, Phys. Lett. A 176, 113 (1993)
7. I. Klaft, S. Borneis, T. Engel, B. Fricke, R. Grieser, G. Huber, T. Kühl, D. Marx, R. Neumann, S. Schröder, P. Seeling, L. Völker, Phys. Rev. Lett. 73, 2425 (1994)
8. W. Quint, J. Dilling, S. Djekic, H. Häffner, N. Hermanspahn, H.-J. Kluge, G. Marx, R. Moore, D. Rodriguez, J. Schönfelder, G. Sikler, T. Valenzuela, J. Verda, C. Weber, G. Werth, Hyperfine Interact. 132, 457 (2001)
9. M. Vogel, W. Quint, Phys. Rep. 490, 1 (2010)
10. E. Riedle, S.H. Ashworth, J.T. Farrell, D.J. Nesbitt, Rev. Sci. Instrum. 65, 42 (1994)
11. T. Führer, D. Stang, T. Walther, Opt. Express 17, 4991 (2009)
12. D.H. McIntyre, T.W. Hänsch, Phys. Rev. A 36, 4115 (1987)
13. G.P. Barwood, W.R.C. Rowley, P. Gill, J.L. Flowers, B.W. Petley, Phys. Rev. A 43, 4783 (1991)
14. P. Cancio, D. Bermejo, J. Opt. Soc. Am. B 14, 1305 (1997)
15. J. Cariou, P. Luc, Atlas du Spectre d'Absorption de la Molécule Tellure (Lab. Aime-Cotton CNRS, Paris, 1980)
16. J.D. Gillaspy, C.J. Sansonetti, J. Opt. Soc. Am. B, Opt. Phys. 8, 2414 (1991)
17. D.H. McIntyre, T.W. Hänsch, Phys. Rev. A 34, 4504 (1986)
18. D.H. McIntyre, W.M. Fairbank Jr., S.A. Lee, T.W. Hänsch, E. Riis, Phys. Rev. A 41, 4632 (1990)
19. B. Edlen, Metrologia 2, 12 (1965)

[^0]:    S. Albrecht • S. Altenburg • C. Siegel $\cdot$ N. Herschbach • G. Birkl ( $\boxtimes$ )

    Institut für Angewandte Physik, Technische Universität Darmstadt, Schlossgartenstraße 7, 64289 Darmstadt, Germany
    e-mail: gerhard.birk1@physik.tu-darmstadt.de

