Fabrication and characterization of super-polished wedged borosilicate nano-cells.

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We report on the fabrication of an all-glass vapor cell with a thickness varying linearly between (exactly) 0 and \( \sim 1 \mu m \). The cell is made in Borofloat glass that allows super polish roughness, a full optical bonding assembling and easy filling with alkali vapors. We detail the challenging manufacture steps and present experimental spectra resulting from off-axis fluorescence and transmission spectroscopy of the Cesium D1 line. The very small surface roughness of 1Å rms is promising to investigate the atom-surface interaction or to minimize parasite stray light.

I. INTRODUCTION

Miniaturization of devices based on atomic vapor has set the stage for the development of various compact light-matter interfaces [1, 2]. However, to reach high precision performances required to be used as atomic sensors [3, 4], integrated optical chips immersed in alkali-vapors demand meticulous characterization. In this perspective, vapor nano-systems [5–7] have recently been used to clarify fundamental problems in optics such as the role of dipole interactions in resonant light scattering [7, 8], control the coherent excitation of Rydberg atoms [9], or evidence the non-local response of an atomic gas [10]. Another issue inherent to the system size reduction, is the growing influence of the long-range atom-surface interaction, that triggered the development of new glass nano-cells [11–13]. Also, by combining short repulsive and long attractive potential, the possibility to trap atoms in bound states has theoretically been predicted [14] and this ability could make way to hybrid nanoscale atom-surface meta-materials [15].

In this article, we report on the fabrication of a glue-free all-glass nano-cell with a thickness varying linearly between (exactly) 0 and \( \sim 1 \mu m \), and with a surface roughness of 1Å rms. Filled with Cesium (Cs), the cell was initially built for the interrogation of small atomic vapor layers of alkali and investigate the hypothetical existence of an atom-surface bound state that would depend on the surface corrugation [16]. Developing surfaces with ultra-low surface roughness is also an asset to reduce parasite scattering limiting signal-to-noise ratio of light matter interaction with nanometric atomic ensembles. Besides, the super-polish surfaces presented here can be ideal candidates for the fabrication of glass micro-cavities for QED experiments [17]. Therefore, the techniques that we present may have a large panel of applications and find audience in diverse domains such as surface science optics and atomic physics. We will first describe the cell fabrication process and how we filled it with Cs. We subsequently present the methods to check the requirements in terms of surface roughness and cell thickness. Finally we bring forward promising examples of spectra in transmission and off-axis fluorescence of the Cs D1 line.

II. FABRICATION AND CHARACTERIZATION

The cell was fabricated at Laboratoire Charles Fabry of Institut d’Optique (Palaiseau, France) and filled with a cesium vapor at the Laboratoire GEPI of Observatoire de Paris (Paris, France). It is made of four parts that are assembled by optical contact bonding leading to a monolithic ensemble (see Fig. 1a). Using a unique material for all parts avoids differential thermal expansion that could damage the optical bonding. Borofloat glass has been chosen for its good optical properties in the visible and near-infrared spectrum, low cost and facility to be super-polished in comparison to other materials used in previous cells (sapphire for instance). This glass was chosen also to facilitate the sealing to the Pyrex side arm that will contain the Cs reservoir, as the thermal expansion coefficients and the softening temperatures of both glasses are similar. However, unlike sapphire, and similar to fused silica, it reacts with alkali at temperature exceeding \( \sim 200°C \). The fabricated cell is therefore a good candidate for operation at moderate temperatures. The central part (A) (see Fig. 1b) is machined using boring-bits so as to let a 6mm external diameter, 4mm internal diameter and 25mm long tube protrude from the front face to allow for connection to a Pyrex loading manifold containing the Cs reservoir. Glass is then removed from the inside of part (A) using milling-bits. In this hollow piece, a thick plate (B), carefully angled on one side is...
FIG. 2. Procedure to realize a thin wedge with thickness varying between (exactly) 0 and 900 nm. Parts (A) and (B) are optically contacted on a parallel plate (P₁). Left: the thickness of part (B) first exceeds that of part (A) by 300 nm. The flat etalon P₂ therefore sits on the ridge of part (B), leading to interference fringes (between part (A) and the flat etalon) with different white light colours on each side of the ridge. Right: after flattening the ridge and equalizing the heights of part (A) and (B), the flat etalon sits equally on parts (A) and (B), leading to equal colour fringes.

introduced. The cell is closed by two thick plates (C_L) and (C_R) wedged by 1° to dismiss unwanted reflections. Plate (B) is optically contacted on one side on plate (C_R). On the other side, the gap formed by the angle between plate (B) and plate (C_L) forms the nano-enclosure and its realization is therefore crucial. The two main manufacturing difficulties that we will detail are: (i) the control of the nano-gap thickness, (ii) the polishing of the different surfaces to reach excellent surface roughness.

The thin wedge of plate (B) is realized by polishing iterations and controls of the wedge thickness and flatness between two polishing steps. This is done interferometrically, using a He-Ne laser and a Fizeau interferometer. The wedge is realized such that 4 fringes appear in the interferogram, parallel to the y axis (see Fig. 1) and with equal interfringe spacing. This corresponds to a thickness variation of 1.2 µm. Also, the absolute thickness is controlled such that, along its thickest edge, the plate thickness exceeds the thickness of plate (A) by ~300 nm (see Fig. 2a), forming a ridge that will eventually be removed. This is controlled mechanically, using an electronic depth gauge with a resolution of 100 nm. Parts A and B are then optically contacted on a parallel plate (P₁) and polished simultaneously so as to remove the 300 nm thick ridge and bring parts (A) and (B) to equal height for closing purpose. This height equalization is controlled interferometrically, using a flat etalon (P₂) as shown in Fig. 2. This procedure ensures that final assembly brings the closing plate (C_L) in optical contact with both parts (A) and (B). The wedge thickness therefore varies between (exactly) 0 and about 900 nm.

To realize surfaces with very low roughness such as those inside the thin wedge, we first grind them finely using alumina abrasives and a brass grinding wheel. The surfaces are then manually polished on a pitched wheel using an aqueous solution of rare earth oxide abrasives with fine particle size (<1 µm). The final polish is performed with an increasingly diluted solution, leading to a super-polish with surface roughness of 1Å rms or less. Figure 3 shows a typical roughness profile, measured using an optical heterodyne profiler (ZYGO 5500) with a sensitivity of 0.2Å rms [18]. The spatial resolution of the profiler is 1 µm, on the order of the distance travelled by the atomic dipoles of the vapor before they reach steady-state (due to collisions inside the gas or radiative decay). To investigate atom-surface interactions over shorter travelling distances and account for the transient response of the vapor to finer details of the surface [10], we characterize the surface on a smaller scale. To do so, we also acquired images of the super-polished surfaces using an atomic force microscope (AFM) in tapping mode, equipped with a Silicium tip (n-type) [19]. The spatial resolution of the AFM imaging is given by the radius of the tip, typically 10 nm or less. Figure 3 compares typical results obtained for surfaces with a super-polish and a standard polish, for which abrasives are rougher and we stopped the polishing procedure before the final dilution step. The roughness is lower for a super-polish than a standard polish, as expected, and it decreases for lesser spatial resolutions as surface height fluctuations are better averaged [20].

All parts of the cell are cleaned with alcohol prior to final assembly by optical contact bonding (no glue is used). This is done at room temperature by pressing lightly onto the parts to be bonded until white light color fringes disappear and are replaced by a uniform dark fringe where the two parts are in contact. The only requirements are the absence of dust and the complementarity of the two surfaces to be bonded: in our case, we found that a flatness difference of 30 nm rms or less was sufficient to
achieve a stress-resistant optical bonding.

Once the cell is assembled we connected it to a turbo-
molecular pumping manifold via a Pyrex loading manifold,
and baked the cell-manifold assembly at 350 – 400°C for
4 days whilst pumping. Cesium was then transferred,
using a low temperature flame, from a cesium ampoule
4 days whilst pumping. Cesium was then transferred,
achieved in situ to maintain the temperature of the wedge
and bake the cell-manifold assembly at 350 – 400°C for
4 days whilst pumping. Cesium was then transferred,
using a low temperature flame, from a cesium ampoule
with breakable seal to the cell reservoir.

We now provide more information about our in situ
measurement of the wedge thickness. We access the lo-
cal absolute thickness $L(x,y)$ using the interferometric
technique described in Ref. [21]. The reflected intensity
$R_w(x)$ resulting from the interference on the thin wedge (see
Fig. 4a) tells about the local thickness: by scanning the
laser position along the $x$-axis of the cell, from the optical
contact to the thickest part of the wedge, we record on a
powermeter the interferometric profile shown in Fig. 4b.
The optical contact is reached for $x \leq x_0$. For $x > x_0$
and a given value of $y$, the data are very well fitted by
the Fabry-Perot model:

$$R_w(x) \propto \frac{F \sin^2(\phi(x)/2)}{1 + F \sin^2(\phi(x)/2)}, \quad \text{(1)}$$

with $F = 4r^2/(1-r^2)^2 = (2F/\pi)^2$ ($F$ is the finesse of
the wedge etalon), $r = (n_g - n)/(n_g + n)$, the reflectiv-
ity (in field amplitude) of the glass-to-vapor interface, $n_g$
the glass refractive index, $n$, the refractive index of the
vapor, and $\phi(x) = 4\pi L(x)/\lambda$, the cumulative phase over
one round-trip of the laser inside the wedge. Here, we
assume that the wedge thickness varies linearly with $x$
for $x > x_0$ and the only free parameter of the fit is there-
for the wedge angle, $\alpha(y) = L(x,y)/(x - x_0)$. Also,
we perform this interferometric mapping of the thickness
at very low vapor pressure and with laser light far de-
tuned from the atomic resonances. The vapor index $n$
can therefore be assimilated to unity. Figure 4c shows
a 2D map of $L(x,y)$. The uncertainty on each point is
$\pm 5$ nm, limited by the finite size of the laser beam (waist
radius $w_0 \approx 40 \mu$m).

III. EXAMPLE OF SPECTRA

Finally we present examples of spectroscopic data ob-
tained with the cell described above. The cell was en-
closed in an oven with small apertures to let a laser
beam propagate through the wedge and to collect the
fluorescence light at 90° from the laser beam. With this
set-up we acquired absorption and fluorescence spectra
shown in Fig. 5. These spectra were obtained by scan-
ing the frequency of a laser diode around the D1 line of
Cs at $\lambda = 894$ nm. In practice one should pay at-
tention to maintain the temperature of the wedge 30°C
higher that the temperature of the Cesium reservoir to
avoid Cesium to condense on the wedge when the tem-
perature is ramped down below the melting point. The

![FIG. 4. (a) Schematic of the experiment. SMFL : single-mode fiber laser; PD : photodiode; APD : avalanche photodiode; NC : nano-cell, PW : power-meter. (b) Reflected interference signal $R_w(x)$ (normalized to the signal obtained at a bright fringe). Black circles : data acquired for $y = 5$ mm. Solid red line : fit by the Fabry-Perot model. For $x < x_0$ (dashed blue rectangle) the wedge thickness is exactly zero since parts (B) and (C1) are optically bonded (see Fig. 1b). (c) 2D-mapping of the wedge thickness (resolution along $x$ and $y : 0.5$ mm).](image1)

![FIG. 5. Spectra of the D1 line of cesium acquired using the nano-cell described in the main text, heated at 174°C. Cell thickness: $L = 447$ nm. (a) In blue circles (red squares) : Normalized fluorescence spectrum collected at 90° from laser propagation (Absorption spectrum). Horizontal axe: laser frequency detuning with respect to the $F = 4 \rightarrow F' = 3$ hyperfine transition of the $6S_{1/2}$ ground state. The peak centered around 1 GHz corresponds to the $F = 4 \rightarrow F' = 4$ transition. The data are binned 10 times by steps of 2 MHz. (b) Fluorescence spectrum from the wedge versus the laser frequency detuning, acquired at 90° from the incoming beam. The data are binned by steps of 2 MHz.](image2)
spectra of Fig. 5 were obtained at a cell temperature of 174°C (temperature of the reservoir, corresponding to an atomic density \( N = 2.6 \times 10^{15} \) cm\(^{-3}\)). The laser was focused onto the thin wedge at a point where the local thickness is \( L = \lambda/2 = 447 \) nm. The laser power was \( \sim 1 \) µW and the laser spot at \( 1/e^2 \) radius was 40 µm. The transmitted light was collected onto a standard Si photodiode. We simultaneously acquired the fluorescence spectrum using a fibred avalanche photodiode in photon counting mode. The overall detection efficiency was \( \sim 1.5 \times 10^{-3} \), accounting for the solid angle collection efficiency (7.5 \( \times 10^{-3} \)), the fiber coupling (50%) and the photodiode quantum efficiency (40%). The possibility to acquire data on a zero background may improve signal-to-noise ratio in comparison with transmission spectra. Both spectra in fluorescence and transmission exhibit a narrow lineshape (smaller than Doppler width), characteristic of the Dicke narrowing observed in nano-cells [?].

IV. CONCLUSION

In conclusion, we have built a new generation of nano-cells that presents many advantages. The cell parts are made of Borofloat glass that allows a very good polishing and therefore a very small surface roughness. It also allows to connect all parts of the cell with optical bonding resulting in a monolithic cell that makes no use of any glue. It therefore avoids any degasing issues for future spectroscopic studies. For the surfaces in contact with the atoms in the nano-wedge, a super-polish is obtained and the surface roughness is less than 1Å rms. The wedge has been realized and characterized using interferometric techniques and the thickness of the cell ranges from exactly zero (optical contact) to 900 nm. Few-nanometer thick atomic vapors have never been probed due to the way nano-cells were closed prior to this work. The type of cells presented here opens up access for the exploration of new mesoscopic systems. The spectra measured using these cells are promising for future investigation of atom-surface interactions. Our fabrication technique allows in principle to cover the cell surfaces with a dielectric layer to increase the reflectivity and hence the dipole-dipole collective effects. The fabrication is theoretically not restricted to glass and testing it with other materials could provide more data on unknown atom-surface potentials. Finally, such cells could be used as in Ref. [23] to perform molecular spectroscopy with various confinement depths.

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