Oscillatory behaviour in Type IA FBG: ruling out chemical complexity

George Simpson, Kyriacos Kalli, John Canning, Amedee Lacraz
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George Simpson a, Kyriacos Kalli * b, John Canning c and Amedee Lacraz b
a Technology Consulting, Deloitte Touche Tohmatsu, Perth, Western Australia
b Nanophotonics Research Laboratory, Cyprus University of Technology, Limassol, Cyprus
c interdisciplinary Photonics Laboratories, School of Chemistry, The University of Sydney, Australia

ABSTRACT

Type IA FBG are regenerated gratings that appear in hydrogenated germanosilicate fibre of all types during prolonged UV exposure. The gratings are characterised by a large Bragg wavelength shift and a concomitant increase in the mean fibre core index. Modulated index changes are complex by comparison and significantly weaker, often characterised by oscillatory growth behaviour. Low thermal stability of Type IA gratings suggests a possible chemical role similar to thermally processed optical fibres where autocatalysis has been observed. We show that GeOH and SiOH formation are not out-of-phase and follow each other, with no evidence of autocatalysis, ruling out a chemical origin.

Keywords: fibre Bragg gratings, optical fibre sensors, absorption spectra, hydrogenation

1. INTRODUCTION

Type IA fibre Bragg gratings (FBG) are formed following the erasure of a Type I grating in hydrogenated germanosilicate fibre of all types at room temperature [1]. They are characterised by a large increase in the Bragg wavelength (ΔλBragg) and a concomitant increase in the mean index (Δn_eff) of the fibre core (Figure 1). Type IA gratings have also been designated Type 1p because the index change is positive [2]. The modulated index change (Δn_mod) is more complicated, and is characterised by oscillatory growth behaviour. The overall grating growth is correlated with absorption features that appear at ~1400nm and that can be deconvoluted into the relevant SiOH and GeOH bands formed in the fibre. The material changes are localised and occur only in the region of UV laser exposure and are initiated by the combination of hydrogenation conditions and UV-laser fluence. This increase in ΔλBragg is dependent on fibre type and the fibre hydrogen loading conditions [3]. In the case of highly doped fibres, such as pure Ge or B/Ge co-doped fibre, ΔλBragg ~ 15–20nm, whereas for hydrogenated SMF28 it is lower at ΔλBragg ~ 5–8nm [3]. Such large values of ΔλBragg translate to an effective core index increase up to Δn_eff ~ 2x10^-2. We believe that these index modifications must reflect material changes within the core and core-cladding regions, consistent with net structural relaxation and therefore stress changes at the core-cladding interface. The extraordinarily large fluence involved further supports an annealing-like relaxation. Moreover, such a large increase in Δn_eff raises the numerical aperture (NA) of the optical fibre so that it becomes few-moded [4] and the NA of the fibre can be increased in excess of 50%. These changes are well beyond the local polarisability changes around altered or induced microscopic defect sites alone. Figure 2 shows the strong correlation between the growth of the OH absorption feature during prolonged UV exposure and the increase in ΔλBragg. The absorption is characterised by the formation of both SiOH and GeOH groups; this becomes more complicated when other fibre dopants are considered. Given that the formation of these species can provide an indication of net energy imparted into the network, high temperature thermal annealing can lead to auto-oscillatory behaviour as hydrogen out-diffuses [5]. The failure to observe this in Type IA grating formation suggests insufficient effective temperatures induced by UV excitation; transient Bragg wavelength shifts during inscription show that it is between 100 and 200°C. We may initially conclude that auto-oscillatory catalysis does not occur and that out-diffusion of hydrogen is relatively low – but is this supposition premature given the role of UV-induced bond breaking? The formation of OH in glass is strongly associated with a net local volume increase leading to relaxation of the tensile stresses at the core-cladding interface. However, volume expansion correlates with a negative index drop convolved with a positive index increase with stress relief leading to a positive index increase. The polarisability of OH has a strong positive index as well. The net result is a large ΔλBragg shift arising from a large dc index change at the core that is not reflected in the index modulation, indicating two different index origins.

* kyriacos.kalli@cut.ac.cy; phone +35725002609; www.cut.ac.cy

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Figure 1. Characteristic red shift during prolonged UV exposure of hydrogenated germanosilicate fibre.

Figure 2. Mean index change (or IA maturity) and the 1400nm absorption band under UV exposure.

So strong is the correlation between OH formation and Δn_eff that by monitoring the growth of the OH absorption (Figure 2), one can determine whether the fibre has “matured” fully, and any gratings written will have the properties of Type IA FBG [4]. This is clear evidence that the dc changes and the grating properties themselves, whilst correlated, are unrelated in origin – the grating is “burnt” into the OH rich glass possibly through a shift of the OH distribution in the highs of the UV interferogram due to bond breaking and diffusion. We note that the peak absorption band loss at ~1400nm is typically ~1.2dB/mm for a B/Ge co-doped fibre. It is clear that there is a complicated process of chemical interaction occurring in the glass matrix that is driven by the presence of hydrogen and the UV-laser fluence. Indeed, it has been noted that autocatalysis and oscillatory reactions can occur in silicate glass systems under certain conditions where diffusion occurs [5]. We can rule this out because the induced temperatures are too low for diffusion between Si and Ge sites. A key result is the large structural change following the substantial formation of SiOH and GeOH sites in the highly Ge-doped fibre, where conditions may differ from those reported previously. Hence we address whether UV irradiation can lower the effective thermal environment required for autocatalysis. To explore this we undertake a study to either observe or rule out the presence of autocatalytic processes during Type IA grating inscription. By monitoring the increase in absorption at 1400nm, with increasing laser fluence, and decoupling the absorption features associated with the presence of SiOH and GeOH, we look for coupling between the two bands in the form of out of phase wavelength oscillations [5]. The logic behind our investigation is summarised here:

1) If oscillations are present and linked to coupling between SiOH and GeOH, their respective absorption bands will appear out of phase, and follow an exponential decay profile because free OH or H will hop between sites and slowly diffuse out of the fibre core. For this to occur, there needs to be equivalence between GeOH and SiOH free energies, which can only be achieved when their free energy of formation is lower than the surrounding energy bath of the network; at room temperature this is impossible and temperatures above 500°C are necessary. Here the effective temperature is ~ 100°C, so we rely on UV bond breaking to reduce the energies of both Ge and Si sites to allow much lower phonon energies to release OH or H, with index linked to differences between the index polarisability at the GeOH and SiOH sites.

2) If we observe that the decoupled absorption features are oscillating in phase then a chemical origin for the observed index modulations is not possible, leading to other directions. Diffusion should also be observed via exponential decay.

2. RESULTS AND DISCUSSION

Gratings were inscribed using the scanning phase mask technique (A_{mask} = 1071.92nm) and the fibre was illuminated with a CW UV Coherent Sabre FreD laser source operating at λ = 244nm. Laser powers of 200mW were delivered to the fibres that were hydrogenated at P = 200 bar pressure and T = 80°C for a 5-day period.

2.1 OH band structure in different fibre types

Gratings were manufactured using the fibre types described in Table 1; 20-mm fibre sections were exposed to blank-beam UV radiation at a scan speed of 1mm/s and were monitored for transmission loss in the ~1400nm region. The OH bands formed in these different fibre types are shown in Figure 3(a). The traces show that B/Ge fibre (blue) and SMF28...
(cyan) both exhibit asymmetric Gaussian absorption peaks. The remaining plots show the OH bands formed in Ge-doped fibres (Verillion and Fibercore), showing two main peaks. All features are predominantly related to the relative contributions of SiOH and GeOH.

<table>
<thead>
<tr>
<th>Fibre reference</th>
<th>Manufacturer</th>
<th>Figure 3a colour convention</th>
<th>Fibre type</th>
<th>NA</th>
<th>Main peak nm</th>
<th>dB/mm</th>
<th>Secondary peak nm</th>
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<td>1406</td>
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Table 1. Summary of the absorption band peaks and dopant levels for the fibres tested. U – Unidentified.

Deconvolution of the PS1250/1500 fibre data shows the presence of two peaks, Figure 3(b). Note that the peak of the curve is just beyond 1400nm under continuous UV exposure, indicating that the anticipated contributions from a mixture of SiOH (1390nm) and GeOH (1410nm) bands are biased in favour of the GeOH bonds for this fibre; this is consistent with both lower free energy of formation for GeOH and the high concentration of Ge in the core. The solid trace represents the absorption band as measured with an optical spectrum analyser (OSA). The dashed and dotted traces represent the de-convolved Gaussian peaks within the absorption band and the bold solid trace shows the addition of these two traces, which closely follows the experimental data. The deconvolved peaks are located at 1401.1nm (attributed to SiOH contribution) and 1424.1nm (attributed to GeOH contribution). However, we note that the deconvolved peaks are shifted from the anticipated peaks at 1390nm (SiOH) and 1410nm (GeOH), which we believe results from the presence of B in the core. There is also no evidence of a hydride band at 1500nm that is most often associated with GeH; this may indicate the rapid conversion of the hydride to the more stable hydroxyl form, an indication of the underlying complexity involved with numerous routes to OH formation. If we monitor the absorption bands of selected fibres under prolonged UV exposure, we observe that there are significant spectral similarities between the different fibre types shown in Figure 4. The B/Ge co-doped fibres (Figure 4(a) and (b)) both show the same fundamental shape as deconvolved in Figure 3(b), whereas the three Ge-doped fibres show the presence of SiOH at 1390nm. By deconvolving the absorption features for the Fibercore fibre (SD079-00B), linked only to SiOH and GeOH bands, we evaluated the band peaks and areas with the laser scan number, Figure 5. It is clear that the curves display some form of oscillation but that it occurs in phase (Figure 5(a)), from which we can exclude case (1) in favour of case (2), suggesting that under our combined optical fibre/inscription conditions autocatalysis does not occur. Figure 5(a) shows the evolution of the peak wavelengths of the SiOH and GeOH bands with increasing exposure, showing an exponential decay profile as H diffuses out of the fibre core (Fibercore SD079-00B). These peak wavelengths asymptote to the anticipated SiOH and GeOH values with each scan, Figure 5(b). Equilibrium is reached less readily for GeOH than for SiOH, which rapidly reaches saturation after 70 scans; this is consistent with there being less SiOH in the core region. Figure 5(c) confirms that there are no out-of-phase oscillations.
Figure 4. Evolution of the OH band associated with the formation of IA gratings during inscription in Verillion IF0100141010 B/Ge (a), Fibercore PS1250/1500 B/Ge (b), Fibercore SD079-00B Ge (c) and Fibercore SD748A-00AB Ge (d). This figure shows that the competing OH species have are weighted according to fibre dopants and doping levels.

Figure 5. (a) Evolution of SiOH and GeOH absorption band peak wavelengths with laser scan number. (b) Superimposed SiOH/GeOH curves to compare growth and equilibrium with laser scan number. (c) Ratio of SiOH/GeOH absorption band peak wavelengths with scan number and their FFT, excluding out-of-phase oscillations.

In conclusion, for Germanium doped fibre (Fibercore SD079-00B), we show that there is no autocatalysis process contributing to Type IA grating formation, despite the role of OH in the large dc shift and possible role as a photorefractive component generating the low thermal stability of the modulated index. This is consistent with a background energy being too low to equalise the energy levels of the general GeO and SiO bond sites for OH formation. The combined UV bond breaking and effective heating are unable to provide this energy. The net dc shift linked to Type IA gratings can effectively be described as part of the continuum of annealing behaviour possible in composite glass fibre systems, especially when H is present, and the role of chemical complexity in elucidating some of the finer details observed during grating growth processes is ruled out for the Type IA grating in this case – i.e. the observed index modulation is not a reflection of change between OH species indicating other sources for this.

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