Recent years have seen tremendous advances in Si photonic systems. The last hurdles to implementation of full Si photonic systems remain the electrically pumped optical amplifiers and lasers in Si using complementary metal-oxide semiconductor (CMOS) compatible technology. The preferred wavelength for such systems is 1.5 μm. Optical gain in Si has been shown recently utilizing the Raman effect and phase matched four wave mixing, although neither is capable of being electrically pumped. One report shows optically pumped lasing in crystalline Si at 1.27 μm, attributed to the so called A center introduced using surface texturing. It is unclear whether this center, whose nature and origin is unknown, can be incorporated using CMOS processes or electrically pumped. The gain curve of this center is too narrow, ~1 nm, for optical amplifiers, where a broad gain spectrum is needed. Here we show gain over 150 nm, centered at 1550 nm, ideally suited for dense wavelength division multiplexing.

Optical gain in semiconductors is almost universally achieved by population inversion of carriers in a p-n junction. This requires high gain as the high doping required leads to high free carrier and Auger losses, and is only achievable in direct gap semiconductors. An alternative is to incorporate a dopant that can be electrically pumped and has internal gain. Although these are often low gain they can be incorporated in to low-loss devices, in principle, enabling net gain.

The rare earth erbium (Er³⁺) has intrinsic gain and is excited electrically in Si via band gap recombination. Although Er has often been mooted as a route to gain in Si, the optical cross sections were supposed too small (~10⁻²⁰ cm²) to achieve sufficient gain, as this would require the incorporation of unrealistic Er concentrations (~10²⁰ cm⁻³). However, if these constraints could be overcome, it would enable the development of fully integrated Si optical amplifiers and lasers.

We report experimental measurements of optical gain due to Er in Si. The low gain of the rare earths and the restricted device dimensions of only a few centimeters in semiconductors have previously prevented the measurement of the gain coefficients. Here we have developed a technique to enable the measurement of low gain systems in short semiconductor waveguides.

Experiments were carried out on samples fabricated in silicon-on-insulator (SOI) material. The high refractive index Si overlayer forms a waveguide between the oxide interface and the surface. The sample and the measurement system are shown in Fig. 1. The sample is mounted inside a cryostat. Photoluminescence (PL) is excited within the waveguide at one end of the sample, by the probe laser, set at 488 nm and modulated at a high frequency. The light passes through a region of the waveguide illuminated by the pump laser and is collected from the far end of the sample. The pump laser, set at 514 nm, is modulated at a much lower frequency and focused to a stripe 250 μm wide and 1 cm long. The output and changes in the output are measured using the technique where two lock-in amplifiers are connected in series. The output intensity is measured as a function of wavelength by the first lock-in amplifier, referenced to the probe frequency. The output of this first lock-in is fed into the second lock-in amplifier. The change, increase or decrease, in the transmitted light is then measured by the second lock-in referenced to the lower modulation frequency. The change in the loss/gain coefficients due to the optical pumping can be calculated directly from the fractional change in the transmitted intensity.
light. The main features of this technique are, firstly, the use of internally generated PL from the sample itself as the probe. This avoids the highly problematic, if not impossible, task of efficiently coupling an external broad band light source into a narrow waveguide at low temperatures down a cryostat. More fundamentally, in short lengths of materials, the change in PL due to the pump laser induced by the pump laser. The curves have been displaced for clarity; in all cases the dashed line is the zero axis.

(b) Data of (a) converted to $\Delta g$, where $\Delta g=[ln(1+\Delta PL/PL)]/L$ with $L$ the pumped length.

FIG. 2. (a) Curve 1 (●) shows the PL due to the probe laser only, taken from the output of lock-in 1. The region between 1500 and 1600 nm has also been expanded vertically to better reveal the structure on the spectrum in this region. Curve 2 (▲), coincident with the origin, is the output from lock-in 2 with the pump laser on and probe laser off. Curve 3 (■) is the change in PL, $\Delta$PL, with both the probe and pump lasers on and gives the change in the PL due to the probe laser induced by the pump laser. The curves have been displaced for clarity; in all cases the dashed line is the zero axis.

Typical luminescence spectra, measured at 100 K and with a pump beam power density of 4 W cm$^{-2}$, are shown in Fig. 2(a). Curve 1 (●) is the PL from the end of the sample due to the probe laser only, measured on lock-in 1. Curve 2 (▲), coincident with the x axis, is the output from lock-in 2 with only the pump laser on. Clearly no PL due to the pump laser is detected showing that lock-in 1 is acting as a perfect blocking filter. Finally, curve 3 (■) shows the change in the transmitted light from the probe due to the pump laser. Curve 1 is typical of the samples. The peaks at 1130 and 1190 nm are the phonon-assisted band-to-band emission from Si and the associated phonon replica. The emission between 1200 and 1500 we believe is associated with residual implant damage. The 1500–1650 region is associated with the Er transitions. The spectral shape for Er$^{3+}$ over this range is typical of that observed previously in Si at this temperature and in particular shows clearly resolved peaks at 1550 and 1595 nm associated with the two main $^4I_{13/2}^-\rightarrow^4I_{15/2}^0$ transitions in the $4f$ shell. It can be seen from the change in PL (curve 3) that the transmitted light has suffered loss at the Si and defect peaks, attributed to the additional absorption due to free carriers generated by the pump laser but shows net material gain where expected in the Er region. Crucially, this result shows that the gain significantly exceeds any losses due to the additional free carriers needed to achieve it. Figure 2(b) shows the fractional change in transmission converted to change $\Delta g$ in the net material gain where $\Delta g=g - \alpha$, $g$ is the gain due to the stimulated emission and $\alpha$ is the optical loss. The losses in regions 1000–1500 nm are clear as is the gain over the Er region from 1550 to 1650 nm. For this particular experimental run and sample, $\Delta g=0.013$ cm$^{-1}$ for the peak gain and the signal to noise ratio is around 15:1, which could be further improved by increasing the averaging time constant on the second lock in. Consequently, we estimate that this technique could detect gain values down to around 0.001 cm$^{-1}$.

We have measured four SOI samples with different Si overlayer thicknesses and background doping to allow a greater range of effective concentrations to be accessed. Samples S1, a 2.1-μm-thick Si heavily doped (2×10$^{17}$ cm$^{-3}$) n-type layer, and S2, a 2.5 μm moderately doped (3×10$^{16}$ cm$^{-3}$) p-type layer, were implanted with Er at five different energies (1.5, 1.0, 0.65, 0.4, and 0.25 MeV) and doses (1.6×10$^{13}$, 1.0×10$^{13}$, 7.5×10$^{12}$, 5.0×10$^{12}$, and 3.3×10$^{12}$ cm$^{-2}$, respectively) to provide an essentially uniform doping of 10$^{18}$ cm$^{-3}$ to a depth of about 0.5 μm. The total implanted dose was 4.18×10$^{13}$ cm$^{-2}$, giving an effective concentration of the dose averaged over the waveguide volume of 2.0×10$^{17}$ cm$^{-3}$ for sample S1 and of 1.7×10$^{17}$ cm$^{-3}$ for sample S2. These samples were subsequently annealed at 850 °C for 1 min in nitrogen ambient. Samples S3 and S4 were phosphorus doped n-type Si layers with doping concentrations of 10$^{14}$ cm$^{-3}$ and 1.5 and 5 μm in thickness, respectively. These were identically implanted with 2×10$^{13}$ Er cm$^{-2}$ at 0.4 MeV and subsequently annealed at 950 °C for 1 min. This corresponds to a peak concentration of 3×10$^{18}$ cm$^{-3}$ and an effective concentration of 1.3×10$^{17}$ cm$^{-3}$ for sample S3 and 4×10$^{16}$ cm$^{-3}$ for sample S4. Prior to the Er implant, samples S3 and S4 were also implanted with 10$^{15}$ B cm$^{-2}$ at 30 keV and annealed at 950 °C for 20 min in nitrogen ambient to form light emitting diode structures. This process has been previously used to improve and in some situations reverse the temperature quenching of the Si and Er luminescence in crystalline Si and provide room temperature operating light emitting diodes.

Figure 3(a) shows $\Delta g$, taken at 1550 nm and normalized to 100 K, as a function of temperature for all the samples. The thermal quenching is significantly reduced in the boron codoped samples and we believe could be completely eliminated in optimized device structures as has been demonstrated.
Terms for cooperative up-conversion and excited state absorption were not included, indicating that these loss mechanisms are negligible in the Er-doped Si system. Values of the maximum change in net gain, $\Delta g_{\text{max}}$, for the differently doped samples are obtained by extrapolation of the fits of $\Delta g$ to the saturation region and are $0.032 \pm 0.001$, $0.0128 \pm 0.0003$, $0.0673 \pm 0.002$, and $0.0267 \pm 0.0007 \text{ cm}^{-1}$ for samples S1, S2, S3, and S4, respectively. Figure 3(c) shows $\Delta g_{\text{max}}$ as a function of the effective Er concentration. For comparison we have also plotted the maximum gain, $g_{\text{max}}^{\prime}$ ($g_{\text{max}}^{\prime} = \sigma N$, where $\sigma$ is the optical emission cross section and $N$ is the effective Er concentration), using the expected value of $1.8 \times 10^{-20} \text{ cm}^2$ for the optical emission cross section. Measured $\Delta g_{\text{max}}$ values are around six times higher in the Er-doped samples and around 30 times higher for the Er samples codoped with boron than expected. This gives an effective lower limit to the optical cross section for the Er in the boron doped samples of $(5.0 \pm 0.1) \times 10^{-19} \text{ cm}^2$. The lower $\Delta g_{\text{max}}$ measurements in the Er only samples is attributed to the faster temperature quenching than in the samples codoped with boron. The mechanism for the increase in the stimulated emission cross section in Si over that seen in glassy hosts is currently unclear. However, a comparable enhancement of gain has been previously observed in Er in SiO$_2$ when Si nanocrystals (nc-Si), which mediate the excitation, have been introduced. In this work they obtained a value for the emission cross section of $(2.0 \pm 0.5) \times 10^{-19} \text{ cm}^2$ for their Er-doped nc-Si sensitized SiO$_2$—half the value obtained here on Er-doped bulk Si. At an Er concentration $10^{19} \text{ cm}^{-3}$ we would expect to obtain a gain of $22 \text{ dB cm}^{-1}$.

In summary we have demonstrated and measured optical gain in Er-doped Si. The values measured far exceed previous expectations of the likely gain values in this system. The gain values obtained mean that the development of optical amplifiers and laser devices using this approach now becomes a realistic possibility at achievable, optically active doping levels. This system therefore offers a potential route to fully CMOS compatible, all silicon, photonic integrated circuits.

FIG. 3. (a) Normalized $\Delta g$ as a function of temperature, measured at 1550 nm, for samples S1 (△), S2 (+), S3 (●), and S4 (□). The solid lines are guides to the eye. Inset shows spectra of $\Delta g$ for sample S2 for three different temperatures: 80 K (solid line), 110 K (dashed line), and 160 K (crossed line). (b) $\Delta g$ measured at 1550 nm for the same samples as a function of pump power density. The solid lines are theoretical fits to the data. (c) A plot of $\Delta g_{\text{max}}$ obtained from the theoretical fits to the data in (b) against the effective Er concentration. The dashed line is $g_{\text{max}}$ calculated using the expected value for the optical cross section of $1.8 \times 10^{-20} \text{ cm}^2$. 

The inset of Fig. 3(a) shows $\Delta g$ spectra for sample S2 at different temperatures. The higher resolution used over the gain region has enabled the expected structure of the two main emission lines of the Er gain curve to be resolved. Figure 3(b) shows the power dependence of $\Delta g$. The solid lines are theoretical fits using the standard rate equation model for the Er$^{3+}$ ion, using only two fitting parameters.