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# Biomedical Microdevices

## Wireless induction coils embedded in diamond for power transfer in medical implants

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# Wireless induction coils embedded in diamond for power transfer in medical implants

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## Abstract

Wireless power and data transfer to medical implants is a research area where improvements in current state-of-the-art technologies are needed owing to the continuing efforts for miniaturization. At present, lithographical patterning of evaporated metals is widely used for miniature coil fabrication. This method produces coils that are limited to low micron or nanometer thicknesses leading to high impedance values and thus limiting their potential quality. In the present work we describe a novel technique, whereby trenches were milled into a diamond substrate and filled with silver active braze alloy, enabling the manufacture of small, high cross-section, low impedance microcoils capable of transferring up to 10 mW of power up to a distance of 6mm. As a substitute for a metallic braze line used for hermetic sealing, a continuous metal loop when placed parallel and close to the coil surface reduced power transfer efficiency by 43%, but not significantly, when placed perpendicular to the microcoil surface. Encapsulation of the coil by growth of a further layer of diamond reduced the quality factor by an average of 38%, which can be largely avoided by prior oxygen plasma treatment. Furthermore, an accelerated ageing test after encapsulation showed that these coils are long lasting. Our results thus collectively highlight the feasibility of fabricating a high-cross section, biocompatible and long lasting miniaturized microcoil that could be used in either a neural recording or neuromuscular stimulation device.

## Key words

Medical implant · Wireless power · Diamond · Braze · Silver ABA · Induction coil

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## 1 Introduction

Medical bionics is the branch of medical science and technology that deals with intractable and incurable medical conditions to reduce their severity or provide therapeutic treatment using prosthetic and neuromodulation devices. The field has significant market potential owing to a wide range of diseases that the technology can target (Cavuoto 2011; Clark 2003; Shepherd et al. 2013). A number of medical implants are very commonly used, such as cochlear implants, intravascular stents, deep brain stimulators and visual prostheses. One notable trend is the increase in the emergence of artificial neural implants that either record from, or electrically stimulate, peripheral or central neurons.

In order to perform this function, implants require a continuous, efficient power supply (Amato et al. 2013) that would preferably transmit power wirelessly, thus minimizing the risk of infection, patient discomfort, pain, health care costs and the need for implantable batteries. Inductive power transfer is therefore a necessary technology in developing safer, more robust devices with lower risk of tissue damage caused by long periods of continuous transcutaneous connections (Amato et al. 2013; Jow and Ghovanloo 2007; Kadefors et al. 1969; Neagu et al. 1997). The efficiency of wireless power-transfer however, strongly depends on a variety of parameters such as, the materials used, coupling and distance between the transfer and receiver coils and their geometrical dimensions, for example the length and cross-section of the coil wire. Besides the outer and inner diameter of the coils, properties of the encapsulation materials influence the overall efficiency of the implant system (Amato et al. 2013). The efficiency of coils is measured by their quality factor (Q-factor), which is defined as the ratio of the electromagnetic energy transmitted ( $E_{tr}$ ) to the receiver coil and energy dissipated ( $E_{dis}$ ) through it according to Equation (1).

$$Q = \frac{E_{tr}}{E_{dis}} = \omega \circ \frac{L}{R_s} \quad \dots (1)$$

Where,  $\omega \circ$  is the operating frequency,  $L$  is inductance and  $R_s$  is series resistance.

In cochlear and retinal implants, while inductive power transfer systems are being used, the receiver coils are large in dimension (in the centimeter range) and implanted subcutaneously behind the ear anchored to the skull (Amato et al. 2013; Jow and Ghovanloo 2007; Li et al. 2011). In recent decades, the development of microelectronics has provided opportunities for the miniaturization of neural implants. Recently, very small devices designed to be implanted within small body structures such as the eye (Ahnood et al. 2015; Ganesan 2014; Ganesan et al. 2014; Garrett et al. 2012; Hadjinicolaou et al. 2012; Lichter et al. 2015b; Maturana et al. 2016; Nayagam et al. 2015; Shivdasani 2013), or inside cortical blood vessels (Oxley et al. 2016) have been developed. The recent surge in interest in electroceuticals for treatment of a range of disorders has led to a myriad of neuromodulation targets, such as the vagus nerve, for regulation of tumour necrosis factor (TNF) associated with autoimmune diseases of the digestive system (Crohn's disease) and rheumatoid arthritis (Martelli et al. 2014; Tracey 2002). Many of these diseases are chronic and neuromodulation devices to treat them or restore sensation (such as cochlear or retinal implants) would ideally be implanted for long time periods. Such devices have the potential to offer significant quality of life improvements to users but a major challenge when devices are small and implanted distant from the skin surface, is the transfer of sufficient power to the implants to enable long term function. For these devices, it is necessary to develop a miniaturised receiver microcoil.

Most commonly, microcoils are fabricated by photolithographic methods. For very fine structures, methods such as X-ray lithography, electron-beam lithography and ion-beam lithography can be employed. Features in the range of 10 nm can be achieved (Broers 1995). Microcoils fabricated using these methods however, are extremely thin in cross-section resulting in high electrical resistance and the inability to deliver sufficient power inductively to the implant. Wen Li *et al.* designed and fabricated a two layered (each layer consisted of 10 turns) gold coil of 3  $\mu\text{m}$  thickness with outer diameter of 9 mm and inner diameter of 5mm on a Parylene C coated silicon substrate via photolithography. The maximum power of about 43 mW could be delivered at a distance of 1 mm using an inductive link but the power was reduced by 62% at 2 mm distance due to divergence of the electromagnetic field and the delivered power reduced to 10mW, where the maximum detectable range was only about 4 mm (Li et al. 2011).

In addition to the coil design and fabrication, the material used for coil wire is also very important. The coil wire material should have high conductivity for efficient power transmission (Amato et al. 2013; Ko et al. 1977;

1 Neagu et al. 1997). Various materials are being used to make coils such as gold (Li et al. 2011), gallium or  
2 gallium alloys and aluminium (Amato et al. 2013; Baj-Rossi et al. 2013). In this study, we considered the use of  
3 silver based active brazing alloy (ABA) for coil fabrication. Silver ABA is currently being used for  
4 interconnections in electronic packaging industry for their solderability (Zhu and Chung 1994) and can also  
5 serve as a metal film conductor having low electrical resistivity of  $2.2 \times 10^{-8}$  ohm.m, only 38% higher than that  
6 of pure silver ( $1.59 \times 10^{-8}$  ohm.m). Silver ABA also produces low porosity films and has excellent film-substrate  
7 adhesion to diamond (Zhu and Chung 1994). These properties make silver ABA an attractive choice of material  
8 to fabricate a coil. However, whilst the material has some excellent properties it is unsuitable for use in medical  
9 implants without encapsulation as it exhibits very poor biocompatibility (Lichter et al. 2015a).

10 Effective power transfer not only requires an efficient inductive link but also requires that losses due to parasitic  
11 inductive currents induced in the encapsulation for instance be minimised. Interactions of the hermetic  
12 encapsulation of the implants with the inductive coils therefore is a design consideration for maximising clinical  
13 effectiveness and safety over the long term (Donaldson 1976; Kirsten et al. 2012; Li et al. 2010). Various  
14 materials have been investigated for encapsulation of medical implants such as parylene C (Amato et al. 2013;  
15 Chen et al. 2008; Li et al. 2011), polyimide (Bongrain et al. 2011) (Lee et al. 2011), rigid titanium boxes,  
16 silicone (Qian et al. 2011), gold ABA (Lichter et al. 2015a) and diamond (Lichter et al. 2015a). Generally,  
17 metals are preferable for hermetic sealing of electronic packages, as polymers can degrade and become porous  
18 while brittle materials such as glasses are at risk of breakage. Metals however, are electrically conductive  
19 effectively forming a 'short-circuited turn' adjacent to the coils when inductive power is transferred to the  
20 implants. This in turn can affect the efficiency of inductive power link due to its eddy current in the magnetic  
21 field (Donaldson 1992) and thus power loss. Thus, it is preferred to encapsulate the coil and electronics package  
22 using a non-conductive material. Note however, even if a non-conductive material can be used for  
23 encapsulation, hermetic sealing may still require a metallic braze line to be incorporated in the device. For  
24 example, Lichter *et al.* employ a Gold ABA braze line as a method of sealing two halves of a diamond hermetic  
25 capsule together (Lichter et al. 2015b) in similar fashion as shown in Fig.1  
26

27 In order to respond to the above limitations, we have developed a new fabrication method of planar square spiral  
28 microcoils made using silver ABA completely embedded within non-conductive diamond. The coils have a  
29 large rectangular cross-section thus reducing resistance and increasing Q-factor. The microcoils in this work are  
30 capable of providing about 10 mW of transferred power across 0.51 k $\Omega$  at a distance of 6 mm which should be  
31 sufficient to power an implanted device for neural stimulation and/or recording. As the toxicity of silver ABA is  
32 an issue, we also describe a method to encapsulate the microcoil with an additional layer of diamond and test its  
33 effects on power efficiency. In addition to this, an accelerated aging test is conducted on the encapsulated  
34 microcoils. Finally, since the work described here extends upon the work of Lichter *et al.*, it was important to  
35 evaluate the impact that an adjacent braze ring might have on inductive power transfer efficiency of a microcoil  
36 embedded in diamond. While a theoretical study for an adjacent metallic braze line is described by Donaldson  
37 N. d. N. (Donaldson 1992), predicting that power transfer efficiency will be reduced due to the braze line  
38 adjacent to the receiver coil, the impact of braze lines on encapsulated coils has not been empirically studied to  
39 our knowledge.  
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## 2 Materials and Methods

### 2.1 Materials

Silver ABA was used for the microcoil containing silver, copper, aluminum and titanium (Ag 92.75%, Cu 5%, Al 1%, Ti 1.25%, Wesgo Ltd.). Gold ABA containing gold, nickel and titanium (Au 96.4%, Ni 3%, Ti 0.3%, Wesgo Ltd.) was utilised as the braze line material. Microcoils were embedded in polycrystalline diamond (PCD) substrates (Element 6 T100 grade). A mixture of sulfuric acid (95-97% reagent grade, ISO) and sodium nitrate mixture was used to clean PCD substrates. Probe connection wires were made of platinum (A-M systems, Pt-Ir, 0.005" bare) and/or copper (Cu) and attached to the coils for measurement purposes. PCD was grown on the exposed surface of microcoils from a mixture of methane and hydrogen gas (BOC Australia) for full encapsulation. Araldite epoxy was used to cover the connection wires during the ageing test.

### 2.2 Design and fabrication method of microcoils embedded in diamond

A flow chart of the design and fabrication process of microcoils is shown in Fig. 2 (a). Planar square spiral coil trenches with different width, depth and separation between two adjacent turns were laser milled into PCD substrates to compare different coil versions. At the two ends of the spiral trench, holes were laser milled right through the PCD substrate for each sample such that the microcoils ended up as pads on the under-side of the substrate during silver brazing for connection of wires. The laser milled PCD substrates were acid boiled in a mixture of 97% sulfuric acid (5 ml) and sodium nitrate (25-50 mg) for one hour. After acid boiling substrates were cleaned sequentially in acetone and distilled water. The width and height of the microcoil trench and the separation between two consecutive coil turns were measured by scanning electron microscopy (SEM; FEI Quanta, FEG 200, Fig. 2b) and optical profilometry (Fig. 2(c)). Silver ABA paste was poured into the coil trench and melted using a MTI vacuum furnace. The furnace was evacuated to a pressure of  $5 - 6 \times 10^{-6}$  Torr and the temperature ramped up to 960 °C. The samples were held at 960 °C for 10 min and then the temperature was ramped down to room temperature gradually. Excess silver ABA was polished out mechanically using a Coborn PL3 Planetary Lapping machine. SEM images were taken post cleaning. Finally connection wires were laser welded onto the pads on the under-side of the microcoils (Fig. 2(d)).

INSERT FIG. 2 HERE

### 2.3 Electrical characterization method

The electrical characterisation of microcoils is the measure of its Q-factor, DC resistance, and inductance. A 30 MHz frequency response analyser (AP Instruments Inc., Model 300) was used for microcoil characterisation. This analyser had a frequency range from 0.10 Hz to 30 MHz. The above electrical parameters were measured with respect to frequency.

### 2.4 Wireless power transfer and braze line effect measurement method

Power transfer and power receiver units were constructed according to the schematic diagram shown in Fig. 3. A power transfer unit was connected to a DC power supply (model TPS-4000). A 3 cm circular transmitter coil (Nucleus patient coil Z209880 manufactured by Cochlear Ltd, Fig. 3) was attached to the power transfer unit. This unit consists of a tuned circuit and integrated RF coil driver. The DC input supply voltage of the power transfer unit was 3.3 volts and input current was approximately 135 mA (135 mV measured across 1  $\Omega$ ). Radio frequency (RF) waves of 4.82 MHz were generated and tuned near to self-resonance of the receiver microcoils by a RF generator (HP Hewlett Packard, 33120A). In order to measure inductive power transfer the microcoils were placed 6 mm away along the axis passing through the centre of the transformer coil. The eight best fabricated microcoils were tested separately using the same receiver unit. A rectifier circuit was used to obtain output power across various loads. Input current and output voltage at different load resistances (R, Fig. 3) were measured with a multimeter (RPG, DM8100). To assess effects of brazing on efficiency, a gold braze line (5.1  $\times$  5.1 mm) fabricated on a separate diamond substrate was introduced either directly on the surface of the

1 microcoil in a parallel orientation (Fig. 3b) or in a perpendicular position 0.2 mm away from the outer diameter  
2 of the microcoils (Fig. 3c) whilst connected to the measurement circuit.  
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## 8 2.5 Coil encapsulation method 9

10 Following initial baseline measurements, an additional layer of PCD was deposited onto exposed surface of the  
11 microcoils in an Iplas microwave plasma-assisted chemical vapour deposition (CVD) system. Microcoils were  
12 seeded before PCD deposition with nanodiamond (Nano Armor) by ultra-sonication in ~3-5 nm  
13 nanodiamond/methanol solution for 5 minutes. Compressed air was used to dry the samples. A gas mixture of  
14 500 sccm hydrogen and 10 sccm methane (BOC Australia, purity 99.99%) was used. During the growth period,  
15 the microwave power was maintained at 900 W, gas pressure at 60 Torr and stage temperature at 800 °C. The  
16 PCD layer was grown for several hours and Elemental analysis (EDS) of the PCD film was performed to  
17 diagnose the exposure of silver through the PCD layer.  
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## 20 2.6 Accelerated ageing method 21

22 Samples were used in an accelerated ageing test. Samples were individually placed in a capped small clean glass  
23 bottle fulfilled with 0.9% medical grade sterile saline (Aerowash Sterile Sodium Chloride Eyewash Solution).  
24 The bottles were transferred to an environmental chamber (Micro Climate Benchtop Test Chamber Cincinnati  
25 Sub-Zero) kept at 85° C. Images were recorded using SEM before and after ageing for time periods of 30 days  
26 corresponding to approximately 30 months of real time equivalent at 37 °C in a saline environment. An aging  
27 factor ( $Q_{10}$ ) of 2 was used for all accelerated aging calculations.  
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## 30 3 Results and Discussion 31

### 32 3.1 Microcoil fabrication outcomes 33

34 Eighteen  $4.6 \times 4.6$  mm microcoils were fabricated using four sets of parameters according to the values presented  
35 in Table-1. Table-1 also provides the measured resistance and Q-factor for each of the four sets of microcoils. In  
36 order to observe the effect of depth and width of microcoils the depth ( $h=100\mu\text{m}$ ) of set 2 and width ( $w=80\mu\text{m}$ )  
37 of set 3 were different from that of the set1 while the outer dimension (4.6 mm) and number of turns ( $N=4$ ) were  
38 same. In case of set 4 the number turns ( $N=20$ ) was more than set 1 ( $N=4$ ) including much wider cross sectional  
39 area of turns. The results show that the Q-factor is dependent on width, depth and number of turns. The actual  
40 values of width, depth and separation between two consecutive turns presented in the table were obtained from  
41 optical profilometry images which were in most cases close to the expected values in parentheses. Generally, the  
42 separation between two consecutive turns was found to be lower than expected while the width of trench was  
43 found to be several micrometers over that of the expected values during laser milling, most likely due to the  
44 conical shape of the laser used for milling. The depth of the coil trench which also varied by several  
45 micrometers in some places was over that of the targeted depth during laser milling. There was difficulty in  
46 obtaining imaging data from the bottom of the trench for microcoil set 2 due to the small width so depth values  
47 for this set were significantly underestimated. As the eight microcoils from set 4 were superior to samples from  
48 the other sets in terms of lower DC resistance and higher Q-factor, in-depth further experiments were conducted  
49 only on this microcoil set.  
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54 Fig. 4 (a) shows an SEM image of one of the microcoils from set 4. The bottom of the coil trench was found to  
55 be narrower than that of the top due to the focusing angle of the cutting laser. A side wall angle was typical  $\leq 11$   
56 degrees in all our samples. SEM images after polishing and cleaning showed instances of small cracks in the  
57 PCD between coil turns as shown in Fig.4 (b). In our study, the cracks did not short two consecutive turns.  
58 However, there is an inherent risk of shorting between turns negatively impacting on Q-factor if these cracks  
59 would fill with braze. These cracks might be reduced by controlling brazing temperature profiles so that PCD  
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substrates do not bend during heating and by using smooth wheel surfaces during mechanical polishing. Also, lower speed polishing may reduce the appearance of cracks.

Table-1: Resistance and Q-factor (average  $\pm$  std. error) of four sets of coils embedded in diamond with several design parameters. Values in parentheses show targeted/expected parameters. \*Difficulty in getting accurate depth data from this set of coils

Sets and number of coils	Outer diameter in mm	Width (w) $\mu\text{m}$	Depth (h) $\mu\text{m}$	Separation Between turns $\mu\text{m}$	Number of turns	Average resistance ( $R_{DC}$ ) $\Omega$	maximum Q-factor
Set 1 (n=4)	4.6	31.20 $\pm$ 1.67 (21)	35.34 $\pm$ 0.78 (35)	62.05 $\pm$ 1.84 (70)	4	5.18 $\pm$ 0.58	1.21 $\pm$ 0.08 at ~11MHz
Set 2 (n=4)	4.6	34.30 $\pm$ 1.85 (21)	39.12 $\pm$ 2.53* (100)	59.80 $\pm$ 1.53 (70)	4	2.38 $\pm$ 0.11	1.98 $\pm$ 0.05 at ~8 MHz
Set 3 (n=2)	4.6	84.39 $\pm$ 1.15 (80)	35.07 $\pm$ 0.17 (35)	70.80 $\pm$ 0.85 (70)	4	1.35 $\pm$ 0.05	2.69 $\pm$ 0.08 at 5.88 MHz
Set 4 (n=8)	4.6	46.3 $\pm$ 0.41 (50)	120.45 $\pm$ 1.15 (140)	54.00 $\pm$ 0.33 (50)	20	2.25 $\pm$ 0.05	3.87 $\pm$ 0.23 at ~3.2 MHz

INSERT FIG. 4 HERE

### 3.2 Electrical characterization

The Q-factors and AC resistance of the microcoils from set-4 are graphically depicted in fig. 5 with respect to frequency. The measured resistance of the microcoils gradually increased as a function of frequency (Fig. 5 (a)). The maximum value of Q-factor of the microcoils was 4.05 at 3.89 MHz (Fig. 5b). The small variation ( $\pm 0.052$ ) in Q-factor and resistance ( $\pm 0.33$ ) provided evidence of the reproducibility in manufacturing as the temperature during brazing was better controlled than that of the other microcoil sets 1-3.

INSERT FIG. 5 HERE

### 3.3 Wireless power transfer and braze line effects

Fig. 6 (a) shows the amount of average power transfer across different load resistances from 0.51 -7.51 k $\Omega$ . The measured maximum output voltage was of 2.41 volts and an average power of 8.64 $\pm$ 1.62 mW (n=6 coils) was obtained across a 0.51 k $\Omega$  load with the highest transmitted power up to 11.38 mW. The power transfer efficiency of the microcoils as a percentage across different load resistances is shown in Fig. 6 (b). The average power transfer efficiency was 1.96  $\pm$  0.35 % across a 0.51 k $\Omega$  load without the presence of any braze line. When comparing our transmitted power results to coils manufactured using lithographic techniques, we were able to transmit almost the same power with a coil of nearly half the dimensions. Also, the coils made using lithographic techniques were not able to send power beyond 4 mm (Li et al. 2011) but our microcoils are capable to transfer power up to 6 mm away along the axis of the transmitter coil. Transmitted power of 10 mW would be sufficient for neural recording by a chip of 32 channels which consumes a total power of 5.4 mW (Wise 2005). This power could also be sufficient for a neuromuscular or optogenetic stimulation device (Lee 2005; Montgomery et al. 2015). Fig. 6 (c) indicates the impact of a braze line on the power efficiency for three of the eight microcoils of higher Q - factors with increasing distance between the braze line and microcoils. It was observed that the effect of a gold braze line oriented parallel to the microcoil did not affect power

1 efficiency when placed more than 2 mm away from the plane of the microcoil (Fig. 6c) but could reduce power  
2 efficiency by up to 43% when placed directly on the coil (0 mm). However, when the braze line was placed  
3 perpendicularly to the plane of the coil 0.2 mm away from the outer edge, the power transfer efficiency was not  
4 affected significantly compared to the power efficiency of the coils without any braze line.  
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### 10 11 12 3.4 Effects of encapsulation and accelerated ageing 13 14 15

16 Encapsulation of microcoils was performed in two stages. Firstly, a group of 4 out of 8 microcoils from set 4  
17 were encapsulated by a PCD layer grown for 5 h applying the methodology described earlier. Secondly a  
18 separate group of 2 microcoils were encapsulated by the PCD layer grown for 14 h with the same conditions as  
19 applied for the first group. The remaining 2 microcoils were subjected to an ageing test without encapsulation.  
20 Microcoils were grouped in a random manner. Fig. 7 (a) and (b) show SEM images at two different  
21 magnifications for one of the encapsulated microcoils. As seen in the images, larger diamond crystals were  
22 found to grow on the space between coil turns while finer crystals grew over the trench. Elemental analysis  
23 (EDS) of the PCD film, revealed no silver peaks for only one of the microcoils encapsulated by PCD using a 5 h  
24 growth time. In contrast, both microcoils in the second group encapsulated by PCD using a 14 h growth time  
25 were free of silver peaks. After electrical measurements on encapsulated microcoils, an ageing test with the  
26 conditions described in methodology was conducted on all samples exhibiting full encapsulation (1 coil from the  
27 5-hour group and both coils from the 14-hour group).  
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39 The ageing test was conducted for 30 days at 85 °C as described in the methods over 3 encapsulated and 2 non-  
40 encapsulated microcoils. Plots depicting the change in Q-factors before and after PCD growth, after an aging  
41 test and eventually after oxygen plasma treatment of the samples are shown in Fig. 7 (c), (d), (e) and (f)  
42 respectively. When performing measurements after encapsulation, it was observed that the resistance of the  
43 microcoils increased and consequently the Q-factor decreased by 38% over the 3 encapsulated samples (Fig. 7  
44 (c), (d) and (e)). The Q - factor of the 5-hour encapsulated microcoil (Fig. 7 (c)) was found to be higher than  
45 that of the two coils in the 14-hour group (Fig. 7 (d) and Fig. 7 (e)), indicating that a thinner PCD layer may  
46 have less impact on the reduction of Q - factor albeit with the high risk of exposure of silver ABA. It is worth  
47 mentioning that the Q - factor improved to some extent after accelerated ageing of the encapsulated samples.  
48 Subsequent resistivity measurements over the grown PCD films showed that the PCD layers on the coils surface  
49 were conductive due the possible presence of graphitic material which may have caused the drop of the Q-  
50 factor. It is important that DC resistance of the microcoils remained the same, which indicates that the  
51 microcoils embedded in diamond were not changed. On the other hand, a nonconductive oxide layer may have  
52 been formed during the ageing period on the PCD, reducing the conductivity of the films to some extent which  
53 improved the Q - factor which led to oxygen plasma treatment on the PCD layer. After oxygen plasma treatment  
54 the Q - factor of PCD encapsulated coils further increased as in Fig. 7 (c, d and e) and PCD film surface became  
55 nonconductive. Thus, overall for future work it will be important to address a fine balance between the risk of  
56 silver exposure and the nonconductive PCD formation by careful periodic control of the encapsulation growth  
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1 time and then oxygen plasma treatment for each layer corresponding to each periodic growth time of PCD on  
2 coil surface.

3 Somewhat surprisingly, there was no significant change in Q - factor before and after accelerated ageing of the  
4 microcoils without encapsulation (Fig. 7 (f)). The most likely explanation for this is that since silver ABA is an  
5 alloy containing silver, copper, aluminum and titanium, it may take more time to degrade compared to when  
6 using a single metal. SEM images were taken after the ageing tests but no visible changes in the diamond  
7 encapsulated samples were observed in case of 14 h growth. There was a tiny spot observed on PCD layer of 5 h  
8 growth film that appeared to be damaged. EDS test was conducted on the PCD films over the samples. EDS test  
9 on PCD films grown for 5 h over microcoils after accelerated ageing did indicate silver at the surface, leading to  
10 the conclusion that 5 h growth time was insufficient to fully encapsulate the coils. PCD layers grown for 14 h  
11 did not exhibit silver peaks during EDS analysis after accelerated ageing. A film thickness of 2-3 $\mu$ m is typical  
12 after 14 h of growth under the reaction conditions used. Full encapsulation of microcoils in diamond is therefore  
13 possible provided the diamond film is grown to a sufficient thickness. Measurement of power transfer efficiency  
14 of fully encapsulated microcoils after ageing tests indicated that up to 2.64 mW power can be transferred over 6  
15 mm of air. While 10mW is ideal, 2.64 mW is still sufficient to power a typical neural recording device or a low  
16 power stimulator (Wattanapanitch and Sarpeshkar 2011).  
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#### 21 **4 Conclusions**

22 We described the fabrication and electrical characterization along with the hermetic encapsulation and longevity  
23 evaluation of microcoils embedded in diamond for wireless power transfer to medical implants. The results  
24 revealed that fabrication of 4.6 mm diameter microcoils, embedded in diamond, was achievable and  
25 reproducible. The microcoils were able to transfer, 10 mW across 0.51 k $\Omega$  at a distance of 6 mm yielding a  
26 power transfer efficiency of approximately 2%. This is sufficient for low powered neural recording and  
27 neuromuscular stimulation. We demonstrated that a continuous short circuit, similar in diameter and close to the  
28 microcoil reduced transfer efficiency significantly. This impacts on devices that are sealed using a continuous  
29 conducting braze line to close the packaging. Microcoils, completely encapsulated in diamond were fabricated  
30 by growing a further layer of diamond over the embedded coils. Growth of the diamond layer over the coils  
31 reduced the coil Q-factor by an average of 38%. Nonconductive PCD may be grown using oxygen plasma  
32 treatment on PCD layers of several nanometers and continuing the PCD growth up to enough thickness (likely  
33 2-3 microns) to encapsulate microcoils which may not change Q - factor. Encapsulated microcoils did not  
34 degrade further during 30 days of accelerated aging at 85 °C in saline (30 months, real time equivalent at 37 °C)  
35 indicating that the diamond encapsulation would be leak free over, at least, 30 months *in vivo*. With the drop in  
36 Q-factor, fully encapsulated microcoils were capable of receiving 2.64 mW of power over a distance of 6 mm  
37 yielding a maximum transfer efficiency of 0.65%. The outer diameter of all microcoils was only 4.6 mm and  
38 therefore the technology is an attractive option for miniaturisation of wireless power delivery to future implants.  
39 As diamond is a hard, hermetic, biopermanent and biocompatible material, these microcoils are promising for  
40 long-term medical implants.  
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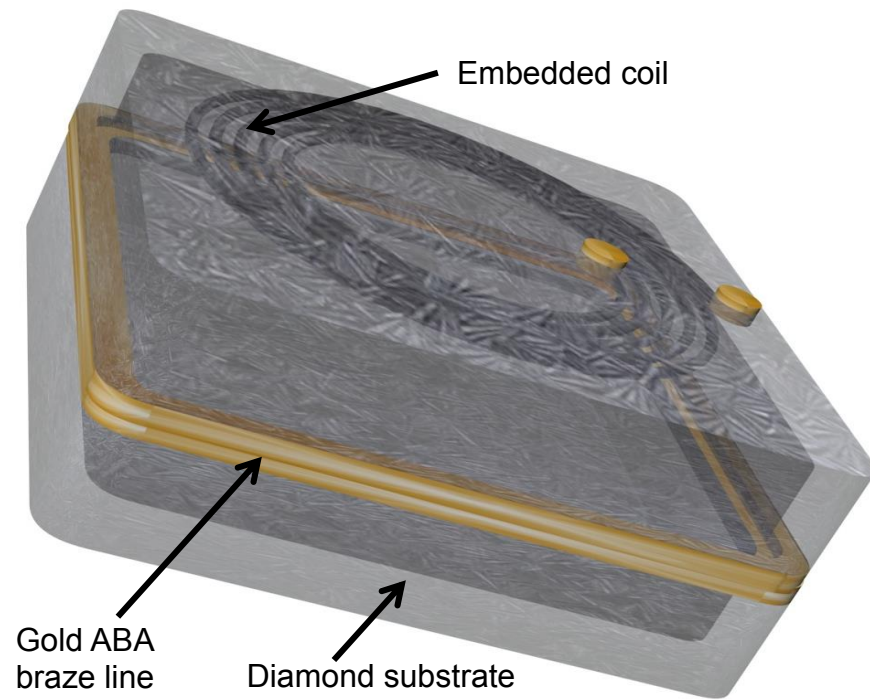
#### 49 **5 Acknowledgement**

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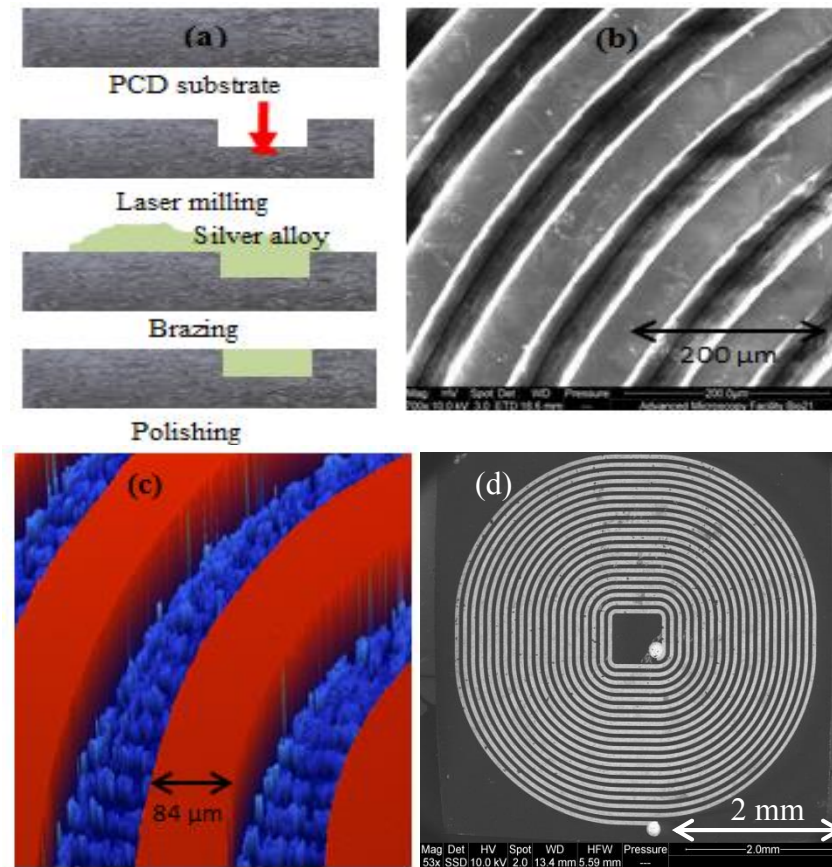
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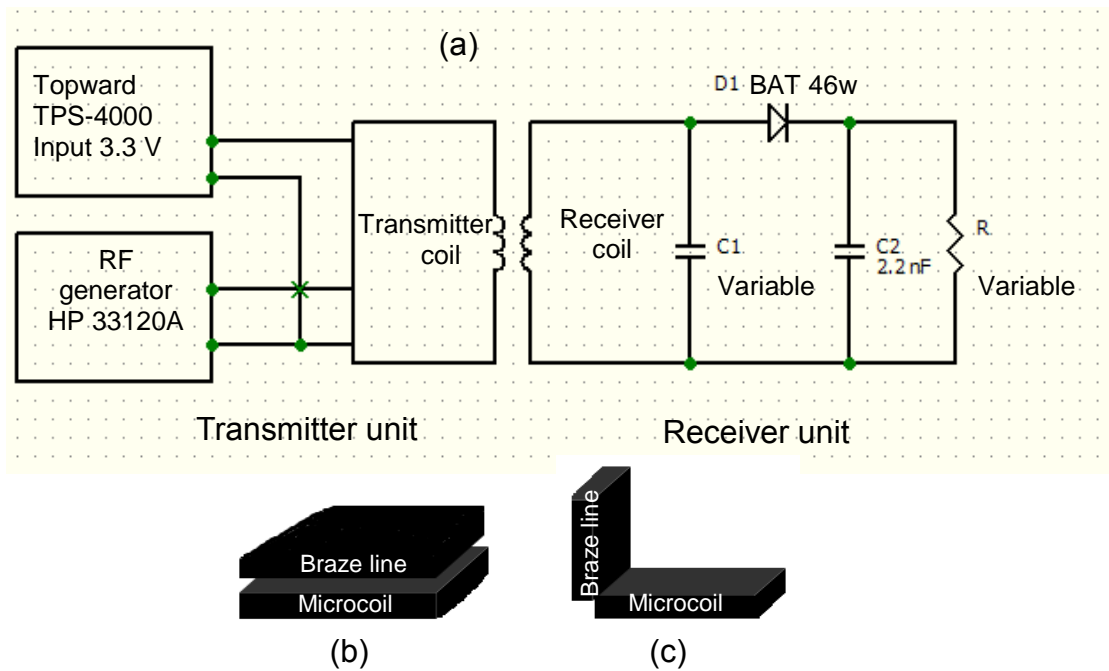
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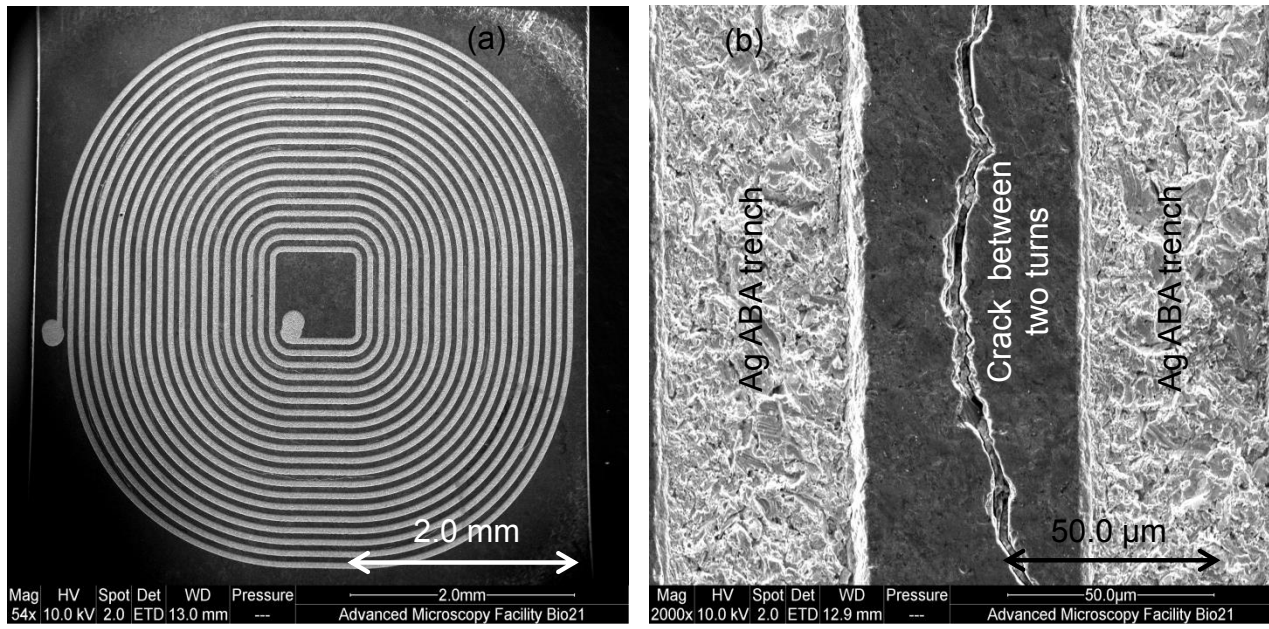
**Fig.1** The schematic diagram of an embedded coil in diamond where the coil is hermetically sealed using laser welding of gold ABA braze lines.



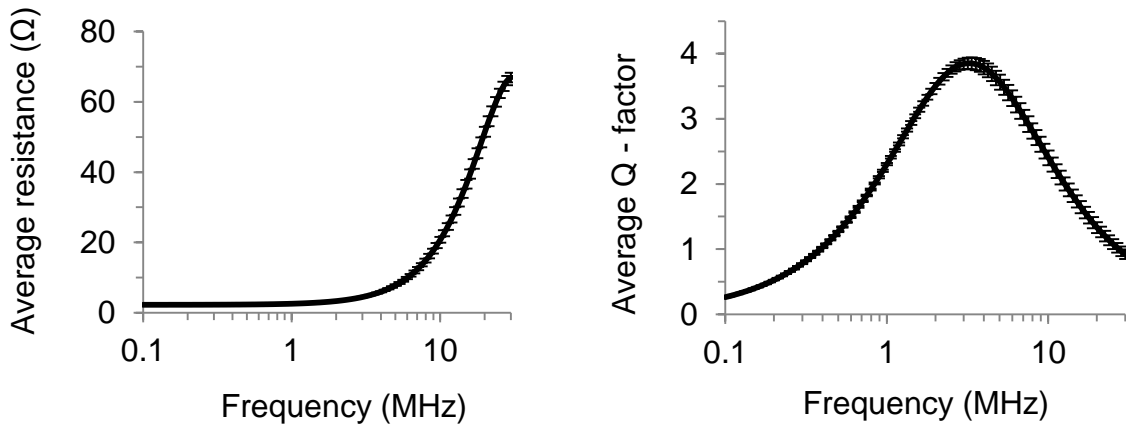
**Fig. 2** (a) Sequence of fabrication process (b) SEM of coil trenches and (c) Optical profilometer image of coil trenches (bottom of the trench is in blue) and (d) Microscopic image of a  $4.6 \times 4.6$  mm coil with 20 turns of silver ABA after fabrication



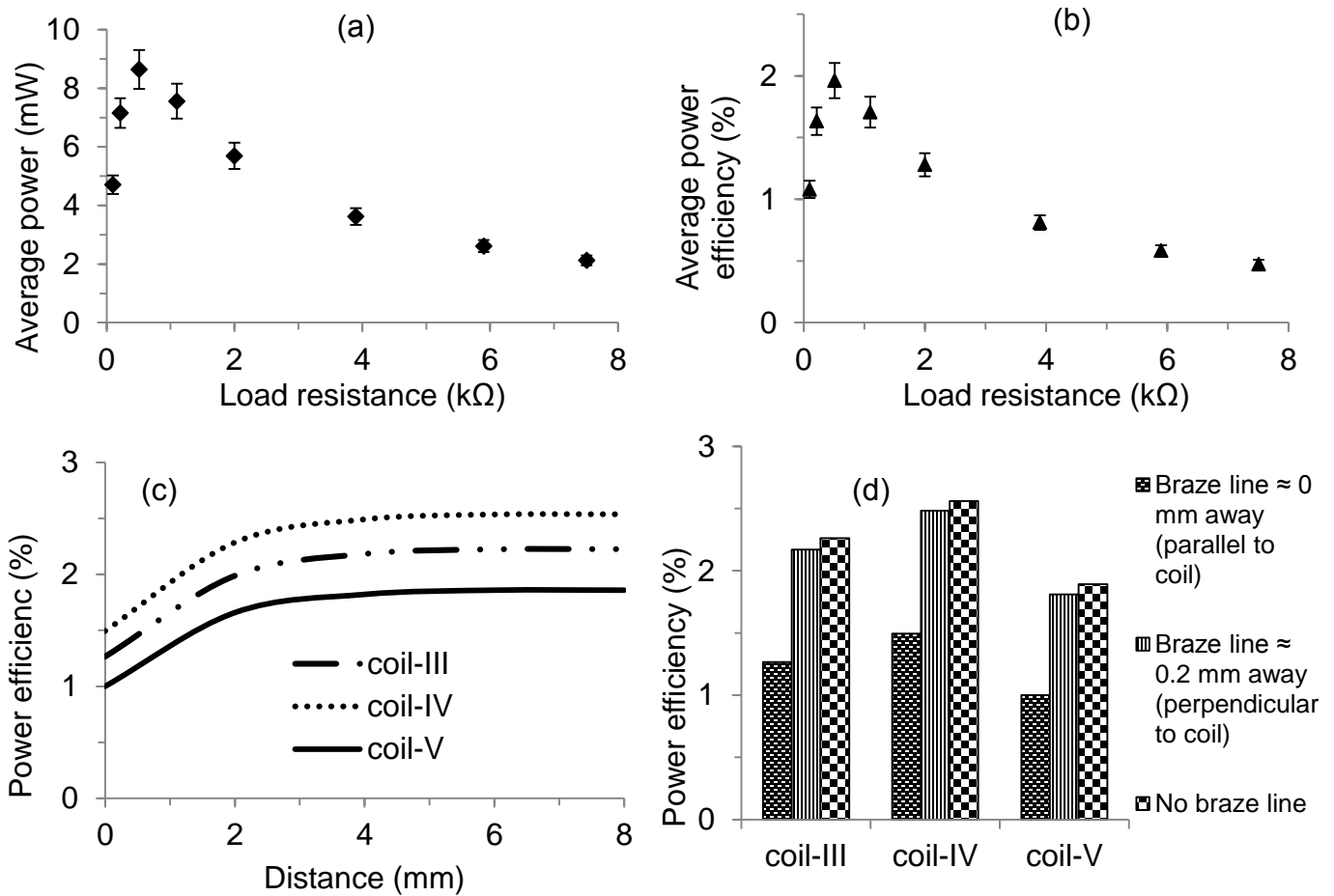
**Fig. 3** Schematic diagrams: (a) circuit of transceiver unit, (b) parallel position of gold braze line to the plane of microcoil and (c) perpendicular position of gold braze line to the plane of the microcoil.



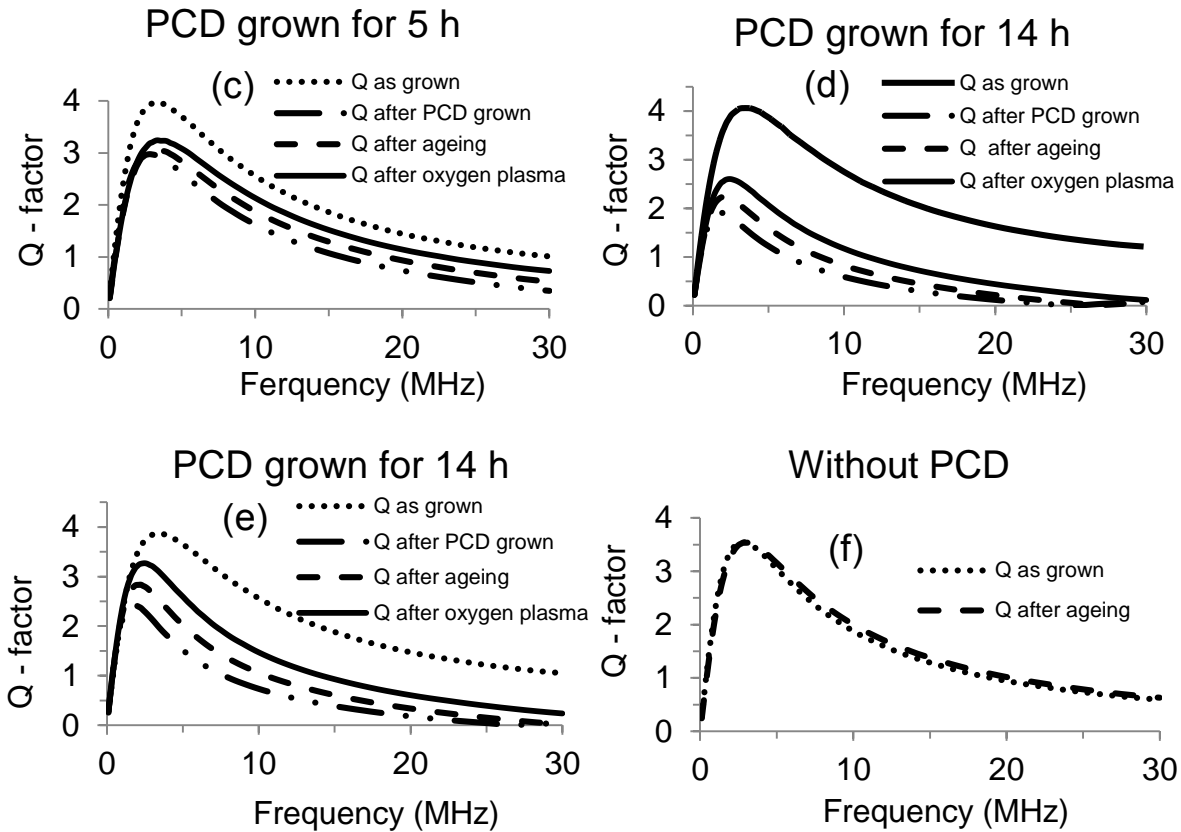
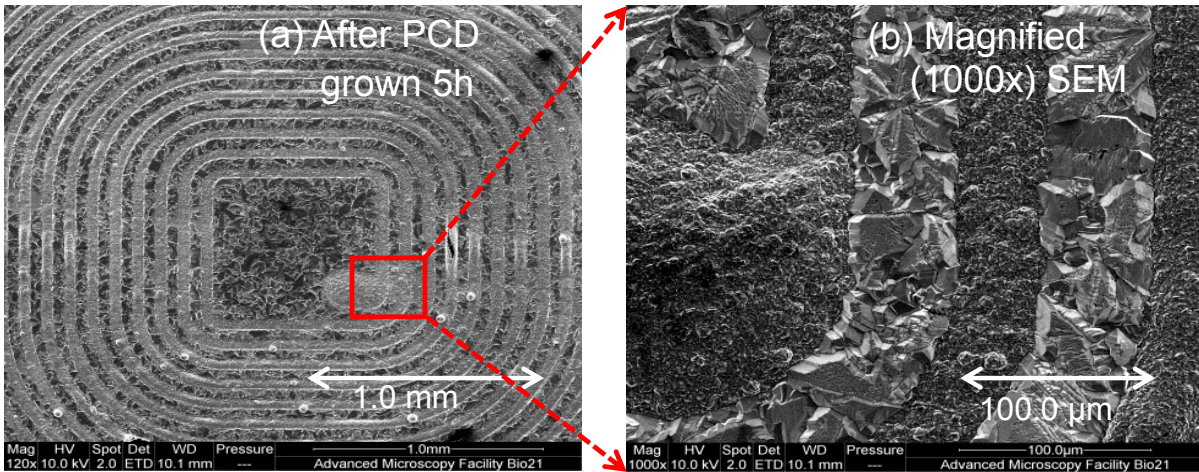
**Fig. 4** (a) Top view SEM image of a microcoil, (b) SEM image of cracks on PCD between two turns



**Fig. 5** (a) Average resistance and (b) Average Q-factor of the eight microcoils from set 4. X-axes are in log scale.



**Fig. 6** (a) Transmitted average power and (b) average power efficiency, plotted against load resistance for the 4.6 mm diameter coils ( $n=6$ ) described in Table 1 from set 4 at a distance of 6 mm between the transmitter and receiver coil. (c) Shows the impact of a parallel gold, braze line ( $5.1 \times 5.1$  mm) on power transfer efficiency of the three highest Q-factor microcoils taken from set 4. The plots show the change in power efficiency (across 0.51 k $\Omega$ ) of the microcoils with increasing distance of the braze line. (d) Shows a comparison of power transfer efficiency for the same three coils as (c) between a braze ring parallel to the coil and a braze line placed 0.2 mm away from the outer edge in a perpendicular orientation with respect to without any braze line.



**Fig. 7** (a) SEM image (top view) of a microcoil after PCD growth on the exposed surface, (b) Magnified (1000x) SEM image of the red box in the Fig. (a). Figures c-f show the change in Q-factor for samples before and after PCD growth, after accelerated aging and after oxygen plasma treatment respectively. The PCD growth times for the samples were (c) 5 h, (d) & (e) 14 h and (f) shows the data corresponding to one of the two samples where the silver coil was aged without PCD growth.