# Time-resolved Density Measurement of Pulse Discharge Plasma by Curling Probe

カーリングプローブによるパルス放電プラズマの時分解密度測定

<u>Anil Pandey</u><sup>1</sup>, Wataru Sakakibara<sup>2</sup>, Hiroyuki Matsuoka<sup>2</sup>, Keiji Nakamura<sup>1</sup> and Hideo Sugai<sup>1</sup> パンデイ アニル<sup>1</sup>, 榊原 渉<sup>2</sup>, 松岡 宏之<sup>2</sup>, 中村 圭二<sup>1</sup>, 菅井 秀郎<sup>1</sup>

> <sup>1</sup>Chubu University, 1200 Matsumoto-cho, Kasugai 487-8501, Japan <sup>2</sup>DOWA Thermotech, 19-1 Ukishima-cho, Mizuho-ku, Nagoya 467-0854, Japan <sup>1</sup>中部大学 〒487-8501 愛知県春日井市松本町1200 <sup>2</sup>DOWAサーモテック〒467-0854 名古屋市瑞穂区浮島町19-1

Curling probe (CP) enables measurement of electron density even in depositive plasmas, measuring a shift of resonance frequency by network analyzer (NWA). In a pulsed discharge, the resonance frequency varies in time, and the frequency spectrum becomes complex and unstable. In order to obtain a stable frequency spectrum during the discharge; the frequency sweep of NWA is triggered by the discharge pulse, in on-sweep mode or on-point mode. The electron density was successfully measured in time-resolved manner for pulsed discharge frequency of 0.4 to 10 kHz.

### **1. Introduction**

A Langmuir probe has been used as a simple plasma diagnostic tool, however it cannot be used in most reactive plasmas since the probe surface is stained with electrically nonconductive layer, thus interrupting the dc current measurement. Recently, a new type of microwave resonator probe, the *curling probe* (CP), has been developed [1] which enables direct measurement of electron density even in depositive plasma. Electron density  $n_e$  (cm<sup>-3</sup>) is obtained from the measured frequency shift of 1/4-wavelength resonance at the frequency  $f_0$  (GHz) in vacuum to  $f_r$  (GHz) in plasma from the following simple equation:

$$n_e = \gamma \frac{f_r^2 - f_0^2}{0.806} \times 10^{10} \quad [\text{cm}^{-3}] \tag{1}$$

Here the coefficient  $\gamma$  is determined by the probe structure and dimensions:  $\gamma = 5.1$  for antenna length of 100 mm and a quartz cover thickness of 0.2 mm.

The CP is compact (minimum diameter ~6 mm) and free from metal impurities owing to alumina or yttria coating. A wide range of electron density  $(10^9-10^{13}\text{cm}^{-3})$  can be measured with high sensitivity. Moreover, an opto-curling probe (OCP) was recently developed [2]. Both CP and OCP are commercially available [3].

To date, the CP has been applied to a constant density *stationary* plasma where the resonance frequency is easily measured by network analyzer (NWA). However, when the CP is inserted into a *pulse-modulated* plasma, the resonance frequency oscillates at the modulation frequency due to temporal change in electron density. As a result, the frequency spectrum becomes complex and varies sweep by sweep of NWA. We have previously reported [4] that a stable frequency spectrum can be obtained with *synchronization* between the pulse period and the NWA sweep period. In this paper, we represent a time-resolved electron density measurement by externally triggering the NWA sweep with *on-sweep mode* or *on-point mode*.

## 2. Experimental

The CP measurement was performed in an apparatus shown in Fig. 1, where a *pulsed* glow plasma is produced in nitrogen at 10 Pa at the pulse frequency 0.4 -10 kHz with duty ratio of 10 - 50 %, applying a negative high-voltage (~ -1.7 kV) to a 12-cm-diam. cathode K in a cylindrical discharge chamber of 60 cm in diameter and 50 cm in length. The CP of 1.6 cm in diameter is set at the center (x=y=z=0) of the chamber, 5 cm above the cathode (y = -5 cm). Five sets of the grounded anode A (30 cm × 30 cm) are installed at the location  $x = \pm 15$  cm, y = 15 cm, y = -17 cm and z = -15 cm.



Fig. 1. Experimental apparatus.

The CP is connected to the NWA (Agilent, model E5071C) which enables time-resolved measurement of reflectance ( $S_{11}$ ), externally triggered by a signal delayed from a synchronous output of the high-voltage pulse generator.

#### 3. On-Sweep Mode Measurement

First of all, the following synchronization condition [4] should be fulfilled to obtain a stable spectrum:

$$(n-1)\frac{T_{\text{pls}}}{T_{\text{swp}}} = m(\text{integer})$$
(1)

where *n* is the number of data points,  $T_{\text{pls}}$  and  $T_{\text{swp}}$  are periods of discharge pulse and frequency sweep, respectively. For example, a pulsed plasma at 0.4 kHz ( $T_{\text{swp}}$ =2.5 ms) and 13% duty ratio was measured as shown in Fig. 2(a) for *m*=10, *n*=1601, and a delay time of 20 µs. This spectrum simultaneously displays 10 spectra at a interval of  $T_{\text{pls}}/m$ =0.25 ms. Analyzing this spectrum, one can obtain a time-resolved reflectance spectrum as shown in Fig. 2(b) where  $\tau$  denotes the time after discharge ignition.



(b) time-resolved data by post data analysis.

The resonance frequency observed in Fig. 2(b) gives the electron density at the discharge time  $\tau$ . In this way, the electron density of 0.4 kHz pulsed discharge (duty ratio 13%) was measured changing the delay time, as shown in Fig. 3.



Fig. 3. Electron density measured by on-sweep mode.

#### 4. On-Point Mode Measurement

For high-frequency discharge pulse (short  $T_{pls}$ ), Eq. (1) requires a short  $T_{swp}$  to get the same time-resolution (same m and n). Then, the frequency sweep is often intermitted during the sweep, which makes post data analysis impossible. In this case, on-point mode is very useful where the NWA digitally sweeps the frequency and measures one data point per each external trigger. Thus, a full spectrum is obtained after n triggers, and one can find the time-resolved spectrum (electron density) by changing the delay time. The measured example is shown in Fig.4 for 1 kHz and 30% duty ratio discharge. The external trigger was delayed by 10µs each, and the resultant time-resolved frequency spectrum gives electron density as a function of discharge time  $\tau$ , as shown in Fig. 5.



Fig. 4. Time-resolved spectrum by on-point mode.



Fig. 5. On-point mode measurement of electron density.

The result of much higher frequency such as 10 kHz will be also reported in the conference.

### References

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