<table>
<thead>
<tr>
<th>Title</th>
<th>Prostate specific antigen density for discriminating prostate cancer from benign prostatic hyperplasia in the gray zone of prostate-specific antigen</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>UNO, Hiromi; KOIDE, Takuya; KURIYAMA, Manabu; BAN, Yoshihito; DEGUCHI, Takashi; KAWADA, Yukimichi</td>
</tr>
<tr>
<td>Citation</td>
<td>泌尿器科紀要 (1999), 45(7): 457-461</td>
</tr>
<tr>
<td>Issue Date</td>
<td>1999-07</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/2433/114086">http://hdl.handle.net/2433/114086</a></td>
</tr>
<tr>
<td>Type</td>
<td>Departmental Bulletin Paper</td>
</tr>
<tr>
<td>Textversion</td>
<td>publisher</td>
</tr>
</tbody>
</table>
PROSTATE SPECIFIC ANTIGEN DENSITY FOR DISCRIMINATING PROSTATE CANCER FROM BENIGN PROSTATIC HYPERPLASIA IN THE GRAY ZONE OF PROSTATE-SPECIFIC ANTIGEN

Hiromi Uno and Takuya Koide
From the Department of Urology, Gifu Prefectural Gero Hot-Spring Hospital

Manabu Kuriyama, Yoshihito Ban, Takashi Deguchi and Yukimichi Kawada
From the Department of Urology, Gifu University School of Medicine

Serum prostate specific antigen (PSA) is currently the best blood marker for prostate cancer. However, low specificity for detection of prostate cancer, especially in the gray zone of PSA, is a problem. We evaluated the clinical significance of PSA density (PSAD) in gray zone PSA cases with conversion of serum PSA to a Stanford reference value.

In a series of histologically confirmed 63 benign prostatic hyperplasia (BPH) patients and 234 prostate cancer patients, 36 BPH patients and 25 prostate cancer patients had gray zone PSA levels. Serum PSA was measured with the Markit-F or Markit-M PA assay. All data were converted to Stanford reference values. We used transabdominal ultrasound to determine prostate volume. PSAD was determined as the serum PSA/prostate volume ratio.

The mean PSA values for BPH and prostate cancer were 6.42±1.80 and 7.80±2.15 ng/ml (p=0.0116), respectively, and prostate volume was 33.4±14.1 ml and 17.1±8.2 ml, respectively (p<0.0001). The mean PSAD for prostate cancer was 0.572±0.363 while that for BPH was 0.218±0.085 (p=0.0001). Cut-off values with sensitivity >90% were 0.218 for PSAD and 30 ml for prostate volume. At these cut-off values, specificity reached 56% for each marker.

In discriminating prostate cancer from BPH in the gray zone of PSA, PSAD demonstrated better performance than PSA.

Key words: Prostate cancer, PSA, PSAD, Cancer screening

INTRODUCTION
Prostate specific antigen (PSA) is a reliable tumor marker in the diagnosis and management of patients with prostate cancer.1,2,3 Assay systems for detecting serum PSA have adequate sensitivity but have not demonstrated sufficient specificity to be used as a screening test in prostate cancer. As PSA is organ-specific but not cancer-specific, above-normal serum PSA levels occur in a high proportion of patients with benign prostatic hyperplasia (BPH). PSA is the least specific in Japanese cases when it ranges from 4.0 to 10.0 ng/ml, the so-called gray zone, in the Hybritech assay because a relatively high percentage of patients with BPH have those values. To enhance the specificity, particularly in the gray zone of PSA, we evaluated PSA density (PSAD) retrospectively. For analyzing data, we converted all serum PSA levels to Stanford reference values to standardize the various PSA assays. Moreover, cases with prostate cancer showing PSAD values less than the cut-off value were examined.

PATIENTS AND METHODS
Histologically confirmed and untreated 63 BPH patients and 234 prostate cancer patients visiting urological hospitals were enrolled for evaluation. Histological diagnosis was performed by random systematic needle biopsy or TUR-P. Serum PSA values were obtained before digital rectal examination using the Markit-F or Markit-M PA assay. All serum data were converted to Stanford reference values according to the literature. Thirty-six BPH patients and 25 prostate cancer patients had gray zone levels of serum PSA, ranging from 4.4 to 11.0 ng/ml (PSA of 4.0 to 10.0 ng/ml in the Hybritech Tandem-R PSA assay was converted to Stanford reference values). Median patient age was 71 years (range 58 to 95) in BPH, and 74 (range 56 to 85) in prostate cancer.

Transabdominal ultrasound was performed using the Aloka SSD-650CL 3.5 MHz, and the entire prostate volume was determined by the following formula: length x height x width x 0.527. The PSAD values were determined as the serum PSA divided by prostate volume.

Student's t test was used to determine statistical significance, and p values <0.05 were regarded as significant.
RESULTS

The pathological grade of the 25 prostate cancer patients with gray zone PSA, was well differentiated in 7, moderately differentiated in 11 and poorly differentiated in 7 patients. Of these patients 72% had clinically localized disease, 5 had stage A, 13 had stage B, 3 had stage C, and 4 had stage D disease.

Serum PSA values converted to Stanford reference values, prostate volume, and PSAD were compared in the overall BPH and prostate cancer groups (Table 1). The mean PSA values for BPH and prostate cancer were 6.4± 1.8 (range ; 4.56 to 10.8) and 7.8± 2.2 (range ; 4.56 to 12.74) ng/ml, respectively, the difference being significant (p=0.0116). Patients with BPH and prostate cancer had a mean prostate volume of 33.4± 14.1 (range ; 13.2 to 78.9), and 17.1 ± 8.2 ml (range ; 5.5 to 42.1), respectively, the difference also being significant (p<0.0001). The mean PSAD for prostate cancer was 0.576±0.363 (range; 0.14 to 1.74) while that for BPH was 0.218±0.085 (range; 0.09 to 0.45) (p=0.0001)

Using various cut-off values of PSAD, we examined the clinical utility. At a PSAD of 0.218 (mean in BPH cases), 92% sensitivity, 56% specificity, and 71% efficacy were obtained (Table 2). In the same manner, we evaluated the significance of prostate volume. There were no cancer cases with a prostate volume of more than 50 ml in our series; sensitivity was 100% at this level. When a prostate volume of 30 ml was used as a cut-off value, the clinical utility was similar to that of PSAD (Table 3). However, the specificity for maintaining sensitivity of about 90% was only 17% with the use of a cut-off value of PSA of 4.8. The 6 prostate cancer patients with the lowest values of PSAD are presented in Table 4. Two of the 25 patients with prostate cancer showed a PSAD of <0.218 and prostate volume of >30 ml (cases 1 and 2). One patient was diagnosed as T2 by digital rectal examination, the other was identified by core needle biopsy because of elevated serum PSA (6.27 ng/ml, T1c).

DISCUSSION

PSA was identified and purified by Wang et al.91 in 1979. Kuriyama et al.101 developed a highly sensitive enzyme immunoassay. PSA is a 34,000 dalton glycoprotein with kinin-kallikrein-like serine protease activity produced by prostatic ductal epithelial cells. PSA has become the most useful tumor marker in the management of prostate cancer, but the lack of specificity for screening of prostate cancer is a problem.11 Diagnosis of prostate cancer is uncommon at a PSA of below 4 ng/ml, and common at that above 10 ng/ml. In the gray zone of PSA between 4.0 and 10.0 ng/ml, the diagnosis of prostate cancer is difficult11. Many tools have been proposed to increase the specificity while maintaining

---

Table 1. Comparison of serum PSA, prostate volume and PSAD in BPH and prostate cancer patients

<table>
<thead>
<tr>
<th>Disease</th>
<th>No. of cases</th>
<th>Age (years)</th>
<th>PSA (ng/ml)</th>
<th>Prostate volume (ml)</th>
<th>PSAD</th>
</tr>
</thead>
<tbody>
<tr>
<td>BPH</td>
<td>36</td>
<td>71±8</td>
<td>6.4±1.8</td>
<td>33.1±14.1</td>
<td>0.218±0.085</td>
</tr>
<tr>
<td>Prostate cancer</td>
<td>25</td>
<td>74±7</td>
<td>7.8±2.2</td>
<td>17.1± 8.2</td>
<td>0.576±0.363</td>
</tr>
</tbody>
</table>

p values

For prostate cancer was 0.576±0.363 (range; 0.14 to 1.74) while that for BPH was 0.218±0.085 (range; 0.09 to 0.45) (p=0.0001)

Table 2. Clinical utility of PSAD according to various cut-off values

<table>
<thead>
<tr>
<th>Cut-off values (level in BPH cases)</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Efficacy</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.218 (mean)</td>
<td>92</td>
<td>56</td>
<td>71</td>
</tr>
<tr>
<td>0.30 (mean+S.D.)</td>
<td>84</td>
<td>86</td>
<td>85</td>
</tr>
<tr>
<td>0.388 (mean+2S.D.)</td>
<td>76</td>
<td>92</td>
<td>85</td>
</tr>
<tr>
<td>0.473 (mean+3S.D.)</td>
<td>56</td>
<td>100</td>
<td>82</td>
</tr>
</tbody>
</table>

Table 3. Clinical utility of prostate volume according to various cut-off values

<table>
<thead>
<tr>
<th>Cut-off values (ml)</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Efficacy</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>72</td>
<td>83</td>
<td>79</td>
</tr>
<tr>
<td>25</td>
<td>88</td>
<td>67</td>
<td>75</td>
</tr>
<tr>
<td>30</td>
<td>92</td>
<td>56</td>
<td>71</td>
</tr>
<tr>
<td>40</td>
<td>96</td>
<td>36</td>
<td>61</td>
</tr>
<tr>
<td>50</td>
<td>100</td>
<td>11</td>
<td>48</td>
</tr>
</tbody>
</table>

Table 4. Prostate cancer cases showing low values of PSAD

<table>
<thead>
<tr>
<th>No.</th>
<th>PSAD</th>
<th>Age (years)</th>
<th>Clinical stage</th>
<th>Pathological grade</th>
<th>PSA (ng/ml)</th>
<th>Prostate volume (ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.14</td>
<td>80</td>
<td>B2</td>
<td>POR</td>
<td>5.70</td>
<td>42.1</td>
</tr>
<tr>
<td>2</td>
<td>0.19</td>
<td>79</td>
<td>B0</td>
<td>MOD</td>
<td>6.27</td>
<td>33.7</td>
</tr>
<tr>
<td>3</td>
<td>0.22</td>
<td>56</td>
<td>B1</td>
<td>WEL</td>
<td>4.75</td>
<td>21.6</td>
</tr>
<tr>
<td>4</td>
<td>0.30</td>
<td>75</td>
<td>A1</td>
<td>WEL</td>
<td>5.89</td>
<td>20.0</td>
</tr>
<tr>
<td>5</td>
<td>0.32</td>
<td>70</td>
<td>B1</td>
<td>WEL</td>
<td>6.76</td>
<td>20.9</td>
</tr>
<tr>
<td>6</td>
<td>0.34</td>
<td>80</td>
<td>D2</td>
<td>WEL</td>
<td>4.76</td>
<td>14.2</td>
</tr>
</tbody>
</table>
sensitivity in the gray zone of PSA. Serum PSA is thought to be dependent on the volume of the prostate and the number of epithelial cells within the prostate. From this finding, Benson et al. introduced the concept of PSAD defined as serum PSA divided by the volume of the prostate, and many studies have been reported. We previously reported the clinical significance of PSAD in distinguishing between BPH and early stages of prostate cancer, when overlapping serum levels of PSA between BPH and early stages of prostate cancer is a problem. Moreover, Benson et al. reported that PSAD offers significant advantages over PSA alone in the evaluation of patients with BPH and prostate cancer in the gray zone of PSA. In their report, the stage of prostate cancer was not mentioned, and they did not discuss prostate cancer cases with values lower than the cut-off values. We have compared the stages and grades of prostate cancer patients and discussed cases showing PSAD values lower than the cut-off values. Fortunately, these cases were diagnosed based upon abnormal digital rectal examination and high PSA value. Although a conclusion cannot be drawn because of the small number of cases examined, the cases with prostate cancer showing low PSAD may be diagnosed by other diagnostic procedures.

The proportion of free PSA to total PSA in the serum is higher in BPH than in prostate cancer. Luderer et al. and Van Cangh et al. have reported that the proportion of free to total PSA ratio (F/T) enhances the ability to distinguish BPH from cancer in the gray zone. However, the advantage of F/T determination is apparent in the higher PSA region of the gray zone. Catalona et al. reported that F/T showed 90% sensitivity and 31.3% specificity in the PSA range of 4–10 ng/ml (Tandem-R PSA). In the range of 2.6–4.0 ng/ml, specificity was only 18% while maintaining the same sensitivity. In Japan, a few clinical studies about the significance of the F/T PSA ratio in the gray zone have been reported and its utility is controversial. Since we did not examine the clinical utility of the F/T PSA ratio, it cannot be compared directly with that of PSAD. However, our PSAD data showed a relatively high clinical utility.

Serum PSA concentration changes in correlation with age, and the rate of change in PSA is greater in men with prostate cancer compared with men without symptoms of prostatic disease. Brawer et al. reported that an annual change in the PSA level of 20% is a significant risk for prostate cancer (PSA velocity). From the point of mass screening, PSA velocity is less useful in the differential diagnosis of BPH and prostate cancer because subjects must be examined annually to determine the need for biopsy. International standardization of PSA has been pointed out because of the various assay kits available. Stamey organized an International Conference for PSA Standardization at Stanford in 1992 and 1994. When analyzing data, calculation of serum PSA values according to the conversion rate for each assay is recommended.

We determined prostate volume using transabdominal ultrasound. Transrectal ultrasound is agreed as the best procedure to detect prostate cancer, but transabdominal ultrasound is thought to be useful in determining prostate volume. Transrectal ultrasound is an important diagnostic procedure, but subjects in a mass screening test may feel inconvenience and discomfort.

CONCLUSIONS

In conclusion, we examined PSA, prostate volume, and PSAD in 36 BPH and 25 prostate cancer patients who had gray zone PSA, and found a significant difference in prostate volume and PSAD. Cut-off values with sensitivity of 92% were 0.218 for PSAD, and 30 ml for prostate volume, and the specificity was 56%. Therefore, the determination of PSAD and prostate volume are useful tools for differential diagnosis of BPH and prostate cancer in the gray zone of PSA. Moreover, if the patients were missed by these strategies, other methods for the diagnosis of prostate cancer such as digital rectal examination, transrectal ultrasound, and total PSA determination may help determine the patients needing a biopsy.

REFERENCES

13) Benson MC, Ring KS and Olsson CA: Determination of stage D-0 carcinoma of the prostate using PSA density. Urologist's Correspondence Club, September 24, 1989
20) van Cangh PJ, De Nayer P, De Vischer L, et al.: Free to total prostate-specific antigen (PSA) ratio improves the discrimination between prostate cancer and benign prostatic hyperplasia (BPH) in the diagnostic gray zone of 1.8 to 10 ng/ml total PSA. Urology 48: 67–70, 1996
24) Catalona WJ, Smith D and Ornstein DK: Prostate cancer detection in men with serum PSA concentration of 2.6 to 4.0 ng/ml and benign prostate examination. enhancement of specificity with free PSA measurement. JAMA 18: 1452–1455, 1997

Received on December 11, 1998
Accepted on May 24, 1999
和文抄録

PSA gray zone の前立腺肥大症と前立腺癌の
鑑別における PSAD の臨床的意義

岐阜県立下呂温泉病院泌尿器科（部長：小出卓也）

宇野 裕巳，小出 卓也

岐阜大学医学部泌尿器科学教室（主任：河田幸道教授）

栗山 学，坂 義人，出口 隆，河田 幸道

前立腺特異抗原（PSA）は特異性が低いため，特に
gray zone における前立腺癌の診断には問題がある。
われわれは PSA 値を Stanford reference value で換算し，gray zone 症例における PSA density (PSAD)
の臨床的意義を検討した。

組織学的診断された前立腺肥大症 (BPH) 63例と
前立腺癌 (PC) 234例のうち，PSA が gray zone を呈
する BPH 36例と PC 25例を対象とした。PSA 値は
Markit-F または MPA で測定し Stanford reference
value を従い換算した。前立腺は経腹的に外腺を含め
て計測し，PSAD 値は PSA 値/前立腺体積で計算した。

PSA 値は BPH で6.42±1.80，PC で7.80±2.15と
なり，BPH との間に p=0.0116 の有意差を認めた。
前立腺体積は BPH で33.4±14.1，PC で17.1±8.2
となり，BPH との間に p<0.0001 の有意差を認め
た。一方 PSAD 値は BPH で0.218±0.085，PC で
0.576±0.363となり BPH との間に p=0.0001 の有
意差を認めた。PSAD および前立腺体積の cut-off 値
を各々の0.218，30ml とした場合，90％以上の感度
を維持しながら特異性はともに56％と良好な結果が得
られた。

Gray zone における PSAD 値の測定は，PSA 単独
に比較して PC と BPH の鑑別診断に有用である可能性を示唆した結果であった。

（泌尿紀要 45：457-461，1999）