

This Provisional PDF corresponds to the article as it appeared upon acceptance. Fully formatted PDF and full text (HTML) versions will be made available soon.

Validation of bidimensional measurement in nasopharyngeal carcinoma

Radiation Oncology 2010, **5**:72 doi:10.1186/1748-717X-5-72

Ting-Shou Chang (aso0225@hotmail.com)
Sau-Tung Chu (enttcd@hotmail.com)
Yu-Yi Hou (hematcd@hotmail.com)
Kuo-Ping Chang (LCC@tzuchi.com.tw)
Chao-Chuan Chi (fangmei2406@hotmail.com)
Ching-Chih Lee (ml2406@hotmail.com)

ISSN 1748-717X

Article type Research

Submission date 11 June 2010

Acceptance date 16 August 2010

Publication date 16 August 2010

Article URL <http://www.ro-journal.com/content/5/1/72>

This peer-reviewed article was published immediately upon acceptance. It can be downloaded, printed and distributed freely for any purposes (see copyright notice below).

Articles in *Radiation Oncology* are listed in PubMed and archived at PubMed Central.

For information about publishing your research in *Radiation Oncology* or any BioMed Central journal, go to

<http://www.ro-journal.com/info/instructions/>

For information about other BioMed Central publications go to

<http://www.biomedcentral.com/>

© 2010 Chang *et al.*, licensee BioMed Central Ltd.

This is an open access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Validation of bidimensional measurement in nasopharyngeal carcinoma

Ting-Shou Chang¹ , Sau-Tung Chu¹, Yu-Yi Hou¹ , Kuo-Ping Chang¹ , Chao-Chuan Chi¹ , Ching-Chih Lee^{2,3*}

¹Department of Otolaryngology, Kaohsiung Veterans General Hospital, Kaohsiung, Taiwan,

²Department of Otolaryngology, Buddhist Tzu Chi Dalin General Hospital,

³School of Medicine, Tzu Chi University, Hualian, Taiwan.

*Corresponding author

Ching-Chih Lee

Department of otolaryngology, Buddhist Tzu Chi General Hospital ; No. 2, Min-Sheng Road, Dalin, Chiayi, Taiwan.

E-mail: ml2406@hotmail.com

Telephone: 886 5 2648000X5231, Fax: 886 5 2648006

E-mail:

Ting-Shou Chang: aso0225@hotmail.com

Sau-Tung Chu: enttcd@hotmail.com

Yu-Yi Hou: hematcd@hotmail.com

Kuo-Ping Chang: LCC@tzuchi.com.tw

Chao-Chuan Chi: fangmei2406@hotmail.com

Ching-Chih Lee: ml2406@hotmail.com

Abstract

Background: Our previous study showed a close relationship between computed tomography (CT)-derived bidimensional measurement of primary tumor and retropharyngeal nodes (BDMprn) and gross tumor volume of primary tumor and retropharyngeal nodes (GTVprn) in nasopharyngeal carcinoma (NPC) and better prognosis for NPC patients with smaller BDMprn. In this study, we report the results on of a study to validate the use of BDM in a separate cohort of NPC patients.

Methods: We retrospectively reviewed 103 newly diagnosed NPC cases who were treated with radiotherapy / concurrent chemoradiotherapy (CCRT) or CCRT with adjuvant chemotherapy from 2002 to 2009. We used magnetic resonance imaging (MRI) to measure BDMprn. We calculated overall survival, recurrence-free and distant metastasis-free survival curves and set a BDMprn cut off point to categorize patients into a high- or low-risk group. We then used Cox proportional hazard model to evaluate the prognostic influence of BDMprn after correcting age, gender and chemotherapy status.

Results: After adjusting for age, gender, and chemotherapy status, BDMprn remained an independent prognostic factor for distant metastasis [Hazard ratio (HR) = 1.046 ; $P = 0.042$] and overall survival (HR = 1.012 ; $P = 0.012$). Patients with $BDMprn < 15 \text{ cm}^2$ had a greater 3-year overall survival rate than those with $BDMprn \geq 15 \text{ cm}^2$ (92.3% vs. 73.7% ; $P = 0.009$). They also had a greater 3-year distant metastasis-free survival (94% vs.75% ; $P = 0.034$).

Conclusion: The predictive ability of BDMprn was validated in a separate NPC cohort. A BDMprn of 15 cm^2 can be used to separate NPC patients into high- and low-risk groups and predict survival rates and metastasis potential. It can, therefore, be used as a reference to design clinical trials, predict prognosis, and make treatment decisions.

Background

Nasopharyngeal carcinoma (NPC) is common among Asians, especially in southern China. While the annual incidence in Western countries is <1 per 100,000 population, it is 6.17 per 100,000 in Taiwan [1]. Because it is difficult to approach nasopharyngeal tumors surgically, chemoradiotherapy or radiotherapy is the primary means of treating this disease [2]. The American Joint Committee of Cancer (AJCC) staging system for NPC is widely used to prognosticate and plan for its treatment and is well-accepted as an evaluation tool in clinical research. However, because the current TNM staging approach is limited in its ability to predict prognosis based on NPC tumor stage [3-4], other factors might be incorporated to further refine prognostic accuracy.

Gross tumor volume is one factor closely related to NPC survival [5-8]. It is not, however, widely advocated as a prognostic factor probably because measuring tumor volume can be time-consuming and labor-intensive. Several studies have used unidimensional and bidimensional measurement to evaluate the tumor size [9-11]. In a previous study, we found bidimensional measurement of primary NPC tumor and retropharyngeal nodes by computed tomography (CT) imaging to be an independent prognostic factor [12]. Due to its improved accuracy, magnetic resonance imaging (MRI) has now virtually replaced CT scan as means of determining the stage of tumors, including NPC, before they are treated [13]. MRI is superior to CT scan for diagnosing the gross extent of tumor infiltration and retropharyngeal lymph node metastasis.

In this study, we retrospectively reviewed MRI images in a separate cohort of NPC patients to further validate of the previous finding regarding the use of bidimensional measurement as means of prognosis in NPC. If confirmed to be an independent prognostic factor, then prognostic ability of the current TNM staging approach can be improved.

Methods

Patient selection

The method of bidimensional measurement of primary tumor and retropharyngeal nodes (BDMprn) in NPC was derived from a cohort of newly diagnosed NPC patients with definite treatment [12]. All patients had histological confirmed NPC and received CT scan of the nasopharyngeal area, chest X-ray, ultrasound or CT scan of the abdomen and whole body bone scan. All cases were restaged based on criteria outlined in the 6th edition of the AJCC staging system [14] Patients received a complete course of radiotherapy (70 Gy - 75 Gy). Patients who received concurrent chemoradiotherapy (CCRT) received three cycles of cisplatin during the same period that were undergoing radiotherapy. Subsequent adjuvant chemotherapy consisting of cisplatin and 5-FU was arranged as guidelines [2]. Using computed tomography-derived measurement, bidimensional measurement of primary tumor and retropharyngeal nodes (BDMprn) in NPC had good correlation with gross tumor volume (Spearman' correlation coefficient=0.845, P<0.001). The intrarater reliability for BDM was good. In multivariate analysis, BDMprn was an independent prognostic factor for any relapse [Hazard ratio (HR)=1.066; P=0.029], and overall survival [HR=1.097; P=0.007]. NPC patients with large BDMprn conferred a poor survival and more recurrences[12].

Validation of the bidimensional measurement of primary tumor and retropharyngeal nodes was performed using a cohort which included NPC patients treated at Kaohsiung Veterans General Hospital from 2002 to 2009. The means of treating NPC patients these two hospitals is similar. All patients received a complete course of radiotherapy (70 Gy - 75 Gy). Concurrent chemotherapy was arranged for NPC patients with advanced T (T2-4) classification or positive neck metastasis. Patients with T2b-T4 or N2-3 disease underwent subsequent adjuvant chemotherapy.

Before treatment, all NPC patients received physical examinations, fiberoptic examinations, chest X-rays, ultrasound or CT scan of the abdomen, whole body bone scan and MRI of nasopharyngeal area. Similarly, all cases were restaged according to the AJCC stage classification system, which was modified in 2002.

MRI technique and measurements

Gross tumor volume of primary tumor and retropharyngeal nodes (GTVprn) of NPC measurement was performed with summation of area technique as previously described [12]. The lateral retropharyngeal nodes were considered malignant if its shortest axial dimension was 5 mm or greater, and any visible node in the median retropharyngeal group was considered metastatic [15-17]. Bidimensional measurement of primary tumor and retropharyngeal nodes (BDMprn) was performed as described previously [12]. Briefly, BDMprn was obtained by multiplying the maximum diameter of the nasopharyngeal tumor and retropharyngeal nodes by the greatest measurement perpendicular to it (Figure 1). Bidimensional measurement of primary tumor (BDMp) was calculated by multiplying the maximum diameter of the nasopharyngeal tumor by the greatest measurement perpendicular to it [18]. It was sometimes difficult to evaluate the anatomic extent of primary tumor and

retropharyngeal node. In such cases, when the outline of tumor was unclear, a radiologist specializing in head and neck cancer helped demarcate the margin. When there was skull base involvement or parapharyngeal space invasion, we could measure the gross tumor and retropharyngeal nodes using the same methods in Figure 1.

The calculation of the three measurements was as the followings:

$GTV_{prn} = \sum \text{Outlined area of primary tumor and retropharyngeal nodes} \times (\text{slice thickness} + \text{split interval}).$

$BDM_{prn} = \sum \text{Maximum diameter} \times \text{greatest perpendicular of primary tumor and retropharyngeal nodes}.$

$BDM_p = \text{Maximum diameter} \times \text{greatest perpendicular of primary tumor}.$

Clinical endpoints

Clinical endpoints were 3-year overall survival, any recurrence and distant metastasis. Six weeks after completing the course of treatment, patients received endoscopy and biopsy of the nasopharynx if necessary. Two months after the course of treatment, each patient received a MRI examination. Chest X-rays, abdominal sonography, and whole body bone scan were performed regularly.

Statistics

Intrarater reliability was measured with using the intraclass correlation coefficient. Overall survival, distant metastasis-free survival and recurrence-free survival were calculated according to the methods of Kaplan and Meier. Differences between multiple survival curves were compared using the log-rank test. The prognostic influence of BDM was assessed using Cox proportional hazards multivariate model after adjusting for age, gender, and chemotherapy status. BDM cut-off values were obtained by receiver operating characteristic (ROC) curve analysis. All statistical

operations were performed using the Statistical Package for Social Sciences, version 15.0 (SPSS, Chicago, IL).

Results

Patient and disease characteristics

The intrarater reliability correlation coefficients for GTVprn, BDMprn, and BDMp were 0.956(0.935-0.97), 0.964 (0.912-0.986), and 0.966 (0.916-0.987). Table 1 shows the characteristics of patients in validation cohort. The mean age was 51±13 years. Of the 103 NPC patients, 77 (75%) patients were men. 88 patients (85%) had advanced stage (stage III-IV). These NPC patients were followed up a median of 43 months (range 9-80 months). Thirty-four (33%) in that cohort had recurrences, including 15 (15%) with locoregional recurrence and 12 (12%) with distant metastasis. Eighteen patients (18%) expired. The 3-year overall survival rate was 87%, locoregional control survival rate 88%, distant metastasis-free survival rate 89%, and recurrence-free survival rate 79%.

Univariate and multivariate analysis

Based on our univariate analysis, bidimensional measurement of primary tumor and retropharyngeal nodes was found to be a significant prognostic factor (Table 2). Adjusting for age, gender, and chemotherapy status, our multivariate analysis found bidimensional measurement of primary tumor and retropharyngeal nodes to significantly predict overall survival (HR=1.012 ; 95% CI: 1.014-1.12 ; P=0.012) and metastasis-free survival (HR=1.046 ; 95% CI: 1.002-1.121 ; P=0.042). The bidimensional measurement of primary tumor was not a significant predictor for outcomes in multivariate analysis. Both univariate and multivariate analysis found

gross tumor volume of primary tumor and retropharyngeal nodes to be a significant prognostic factor.

Bidimensional measurement and risk groups

We wanted to further validate the prognostic ability of bidimensional measurement of primary tumor and retropharyngeal nodes using MRI findings. After analyzing trade-off, we chose 15 cm² as the cut-off point in the validation cohort (additional file 1). Using this cut-off point, we further divided validation cohort into a smaller BDMprn group (67%) and a larger BDMprn group (33%). The smaller BDMprn group had greater 3-year overall survival, distant metastasis-free survival, and recurrence-free survival rates than the large BDM group. (92.3% vs. 73.7% , P =0.009 ; 94% vs. 75%, P=0.034 ; 64.1% vs. 59.7%, P =0.082) (Figure. 2A and 2B), and they were at lower risk.

Discussion

In a previous study, CT-derived bidimensional measurement of primary tumor and retropharyngeal nodes could be used to predict prognosis of NPC [12]. Using MRI to validate the ability of bidimensional measurement of primary tumor and retropharyngeal nodes to predict NPC outcomes in a validation cohort, we found BDMprn remained an independent prognostic factor for overall survival as well as metastasis-free survival. Adopting a BDMprn of 15 cm² as cut-off point in validation cohort, NPC patients whose BDMprn was less than 15 cm² had a better 3-year overall survival rate and distant metastasis-free survival rate than those with BDMprn above this cut off point. Based on these two studies, we have found that BDMprn can be

used to stratify patients into two different prognostic groups with significantly different overall survival and metastatic rates.

Although the current TNM staging system for NPC is widely used, it has been reported to have several deficiencies. Mao *et al.* [3] and Cheng *et al.* [4] have not found any significant differences in local-relapse free survival among the T1, T2, and T3 NPC subgroups. Recently, gross tumor volume has been reported to be a risk factor for local recurrence of NPC [5, 6,19]. However, measurement of gross tumour volume is time-consuming, and the technology, expertise, and manpower are often not available in routine clinical practice.

In a study of bidimensional and unidimensional MRI-derived measurement to reflect NPC tumor anatomic extent at diagnosis or the change in size after treatment, King *et al.* [18] found that BDM of primary tumor was a quicker and more widely applicable method than tumor volume measurement and that it could be used to assess tumor response. However, measurement of retropharyngeal nodes were not included in that series. Tang *et al.* [20] showed that retropharyngeal lymph node metastasis affects the distant metastasis-free survival rates of NPC, and Wang *et al.* [21] found a good correlation between retropharyngeal lymph node metastasis and parapharyngeal space involvement as well as metastasis to Level II, III, IV and/or V nodes. Based on these findings, our previous study modified the approach used by King *et al.* to include both primary tumor and retropharyngeal lymph nodes measurements in our definition of BDM. Previous study found a very close relationship between CT-derived BDMprn, gross tumor volume of primary tumor and retropharyngeal nodes, and overall survival [12]. In the present study, also incorporating retropharyngeal lymph node

measurements, we found MRI-derived BDMprn could also predict overall survival as well as metastasis free survival in NPC patients.

In our study, we found that we could use BDMprn to categorize patients into low- and high-risk groups. This distinction would facilitate treatment decisions, as it would spare low-risk NPC patients from receiving aggressive treatment. Although NPC is markedly radiosensitive, there is a high failure rate in treatment due to its metastatic behavior. Improvement in the outcome for NPC relies on the delivery of higher radiation doses [22]. While radiotherapy is the only standard treatment for early-stage NPC (stage I), the combination of cisplatin-based chemotherapy and radiotherapy is used to treat patients with advanced NPC (stage II-IVB) [2]. The latter group not only receives higher doses of radiotherapy, they also receive chemotherapy, both associated with significant comorbidity, including myocardial infarction, severe nutrient deficiency, nephrotoxicity, transverse myelitis, leukopenia, and central nervous system disease [2, 23-24]. Recent study revealed that NPC patients with $GTV_{prn} \geq 13$ ml conferred a poor prognosis and may benefit from ≥ 4 cycles of chemotherapy [25]. This series implied that high-risk NPC patients, such as large GTV_{prn} , could benefit from more intensive chemotherapy and radiotherapy. The treatment goals for NPC is to adjust chemoradiotherapy dosages to achieve adequate anticancer effects without overly increasing the development of such complications. It would be important and valuable if high-risk NPC patients could be identified in order to adjust the radiation dose and tailor chemotherapy protocol. In this way, high-risk patients (larger BDMprn) may benefit from more extensive treatment approaches, such as more intensive chemotherapy or higher dose of radiation, whereas low-risk

patients (smaller BDMprn) may do well with standard therapy and can be spared the severe toxic side effects of more radical therapy.

Conclusion

We have validated bidimensional measurement of primary tumor and retropharyngeal nodes in a different cohort of NPC with pretreatment staging by MRI. BDMprn, derived by MRI, is closely related to survival rates and metastatic rates of NPC patients. BDMprn can stratify patients into two different prognostic groups with significantly different overall survival. Nasopharyngeal carcinoma patients with large bidimensional measurement have poor survival rates and high metastasis potential. BDMprn might be used in the future for the design of clinical trials, the prediction of survival, and treatment decisions.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

TSC and CCL designed the study, collected the data, interpreted the results of the study, and oversaw the project completion. STC, YYH, KPC and CCC participated in preparing of data acquisition. TSC and CCL performed the statistical analysis and drafted the manuscript. All authors contributed to the scientific setup of the study and revised the manuscript critically, and they have approved the final version of the manuscript.

References

1. Department of Health, The Executive Yuan: **Cancer registry annual report, Republic of China, 2001.** 2004.
2. Al-Sarraf M, LeBlanc M, Giri PG, Fu KK, Cooper J, Vuong T, Forastiere AA, Adams G, Sakr WA, Schuller DE, Ensley JF: **Chemoradiotherapy versus radiotherapy in patients with advanced nasopharyngeal cancer: phase III randomized Intergroup study 0099.** *J Clin Oncol* 1998, **16**:1310-1317.
3. Mao YP, Xie FY, Liu LZ, Sun Y, Li L, Tang LL, Liao XB, Xu HY, Chen L, Lai SZ, et al: **Re-evaluation of 6th edition of AJCC staging system for nasopharyngeal carcinoma and proposed improvement based on magnetic resonance imaging.** *Int J Radiat Oncol Biol Phys* 2009, **73**:1326-1334.
4. Cheng SH, Tsai SY, Horng CF, Yen KL, Jian JJ, Chan KY, Lin CY, Terng SD, Tsou MH, Chu NM, et al: **A prognostic scoring system for locoregional control in nasopharyngeal carcinoma following conformal radiotherapy.** *Int J Radiat Oncol Biol Phys* 2006, **66**:992-1003.
5. Sze WM, Lee AW, Yau TK, Yeung RM, Lau KY, Leung SK, Hung AW, Lee MC, Chappell R, Chan K: **Primary tumor volume of nasopharyngeal carcinoma: prognostic significance for local control.** *Int J Radiat Oncol Biol Phys* 2004, **59**:21-27.
6. Chu ST, Wu PH, Chou P, Lee CC: **Primary tumor volume of nasopharyngeal carcinoma: prognostic significance for recurrence and survival rate.** *Eur Arch Otorhinolaryngol* 2008, **265 Suppl 1**:S115-120.

7. Chen MK, Chen TH, Liu JP, Chang CC, Chie WC: **Better prediction of prognosis for patients with nasopharyngeal carcinoma using primary tumor volume.** *Cancer* 2004, **100**:2160-2166.
8. Chua DT, Sham JS, Kwong DL, Tai KS, Wu PM, Lo M, Yung A, Choy D, Leong L: **Volumetric analysis of tumor extent in nasopharyngeal carcinoma and correlation with treatment outcome.** *Int J Radiat Oncol Biol Phys* 1997, **39**:711-719.
9. World Health Organization.: *WHO handbook for reporting results of cancer treatment.* Geneva Albany, N.Y.: World Health Organization ; sold by WHO Publications Centre USA; 1979.
10. Miller AB, Hoogstraten B, Staquet M, Winkler A: **Reporting results of cancer treatment.** *Cancer* 1981, **47**:207-214.
11. Therasse P, Arbuck SG, Eisenhauer EA, Wanders J, Kaplan RS, Rubinstein L, Verweij J, Van Glabbeke M, van Oosterom AT, Christian MC, Gwyther SG: **New guidelines to evaluate the response to treatment in solid tumors. European Organization for Research and Treatment of Cancer, National Cancer Institute of the United States, National Cancer Institute of Canada.** *J Natl Cancer Inst* 2000, **92**:205-216.
12. Lee CC, Ho HC, Su YC, Lee MS, Hsiao SH, Hwang JH, Hung SK, Chou P: **Bidimensional measurement of nasopharyngeal carcinoma: a simple method to predict outcomes.** *Clin Otolaryngol* 2009, **34**:26-33.
13. Chung NN, Ting LL, Hsu WC, Lui LT, Wang PM: **Impact of magnetic resonance imaging versus CT on nasopharyngeal carcinoma: primary tumor target delineation for radiotherapy.** *Head Neck* 2004, **26**:241-246.
14. *AJCC cancer staging handbook.* New York: Springer; 2002.

15. Ichimura K.: **Can Rouviere's lymph nodes in non-malignant subjects be identified with MRI?** *Auris Nasus Larynx* 1993;**20**; 117-123.
16. Watarai J, Seino Y, Kobayashi M, Shindo M, Kato T: **CT of retropharyngeal lymph node metastasis from maxillary carcinoma.** *Acta Radiol* 1993;34 492-495.
17. King AD, Ahuja AT, Leung SF, Lam WW, Teo P, Chan YL, Metreweli C: **Neck node metastases from nasopharyngeal carcinoma: MR imaging of patterns of disease.** *Head Neck* 2000;**22**; 275-281.
18. King AD, Zee B, Yuen EH, Leung SF, Yeung DK, Ma BB, Wong JK, Kam MK, Ahuja AT, Chan AT: **Nasopharyngeal cancers: which method should be used to measure these irregularly shaped tumors on cross-sectional imaging?** *Int J Radiat Oncol Biol Phys* 2007, **69**:148-154.
19. Shen C, Lu JJ, Gu Y, Zhu G, Hu C, He S: **Prognostic impact of primary tumor volume in patients with nasopharyngeal carcinoma treated by definitive radiation therapy.** *Laryngoscope* 2008, **118**:1206-1210.
20. Tang L, Li L, Mao Y, Liu L, Liang S, Chen Y, Sun Y, Liao X, Tian L, Lin A, et al: **Retropharyngeal lymph node metastasis in nasopharyngeal carcinoma detected by magnetic resonance imaging : prognostic value and staging categories.** *Cancer* 2008, **113**:347-354.
21. Wang XS, Hu CS, Ying HM, Zhou ZR, Ding JH, Feng Y: **Patterns of retropharyngeal node metastasis in nasopharyngeal carcinoma.** *Int J Radiat Oncol Biol Phys* 2009, **73**:194-201.
22. Perez CA, Devineni VR, Marcial-Vega V, Marks JE, Simpson JR, Kucik N: **Carcinoma of the nasopharynx: factors affecting prognosis.** *Int J Radiat Oncol Biol Phys* 1992, **23**:271-280.

23. Erkal HS, Serin M, Cakmak A: **Nasopharyngeal carcinomas: analysis of patient, tumor and treatment characteristics determining outcome.** *Radiother Oncol* 2001, **61**:247-256.
24. Teo PM, Leung SF, Chan AT, Leung TW, Choi PH, Kwan WH, Lee WY, Chau RM, Yu PK, Johnson PJ: **Final report of a randomized trial on altered-fractionated radiotherapy in nasopharyngeal carcinoma prematurely terminated by significant increase in neurologic complications.** *Int J Radiat Oncol Biol Phys* 2000, **48**:1311-1322.
25. Lee CC, Huang TT, Lee MS, Hsiao SH, Lin HY, Su YC, Hsu FC, Hung SK: **Clinical application of tumor volume in advanced nasopharyngeal carcinoma to predict outcome.** *Radiat Oncol* 2010;5:20.

Table 1 Patient Characteristics

Variables	Validation cohort (<i>n</i> =103) No. of patients (%)
Age (years)	
Mean±SD	51±13
Gender	
Male	77(75)
Female	26(25)
Stage	
I	3(3)
II	12(12)
III	56(54)
IV	32(31)
T classification	
T1	25(24)
T2	20(19)
T3	34(33)
T4	24(23)
N classification	
N0	6(6)
N1	11(11)
N2	72(70)
N3	14(14)
Histology grade	
Non-keratinising carcinoma	13(13)
Undifferentiated carcinoma	90(90)
Treatment modality	
RT/CCRT	76(74)
CCRT+CT	27(26)

SD, standard deviation; RT, radiotherapy; CCRT, concurrent chemoradiotherapy; CCRT+CT, concurrent chemoradiotherapy with adjuvant chemotherapy

Table 2 Univariate and multivariate analysis results (n=103)

	Overall survival		Metastasis		Any recurrence	
	Univariate HR (95% CI)	Multivariate* HR (95% CI)	Univariate HR (95% CI)	Multivariate* HR (95% CI)	Univariate HR (95% CI)	Multivariate* HR (95% CI)
GTVprn	1.07 (1.037-1.103) P<0.001	1.069 (1.033-1.107) P<0.001	1.049 (1.013-1.085) P=0.007	1.05 (1.01-1.091) P=0.013	1.048 (1.018-1.079) P=0.002	1.037 (1.004-1.071) P=0.028
BDMprn	1.071 (1.021-1.122) P=0.004	1.012 (1.014-1.12) P=0.012	1.06(1.006-1.116) P=0.028	1.06 (1.002-1.121) P=0.042	1.046 (1.002-1.093) P=0.041	1.038 (0.99-1.088) P=0.121
BDMp	1.056 (1.001-1.113) P=0.045	1.048 (0.991-1.108) P=0.099	1.041 (0.983-1.116) P=0.169	1.043 (0.981-1.108) P=0.176	1.043 (0.993-1.096) P=0.091	1.037 (0.984-1.092) P=0.173

GTVprn, gross tumor volume of primary tumor and retropharyngeal nodes; BDMprn, bidimensional measurement of primary tumor and retropharyngeal nodes; BDMp, bidimensional measurement of primary tumor ;HR, hazard ratio; 95% CI, 95% confidence interval.

*Multivariate analysis: adjusted for age, gender, and chemotherapy status.

Figure legends:

Figure 1

T2-weighted postcontrast MR image in the axial plane. The bidimensional measurement of primary tumor and retropharyngeal nodes (BDMprn) was obtained by summation of multiplying the maximum diameter by the greatest measurement perpendicular to it in nasopharyngeal tumor (A) and retropharyngeal nodes (B). $BDMprn (cm^2) = A1 \times B1 + A2 \times B2 + A3 \times B3$.

Figure 2

Survival curves.

(A)Probability of overall survival rates by small versus large BDMprn. (B)

Probability of distant metastasis-free survival rates by BDMprn.

Additional files:

Additional file 1: Table S1

Validity of BDMprn using death, distant metastasis or any recurrence as the standard.

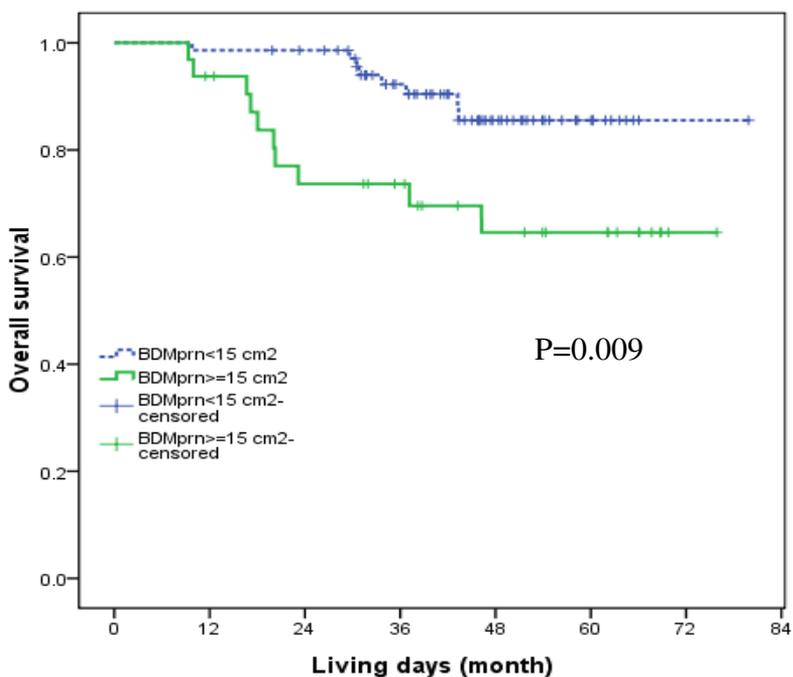


A

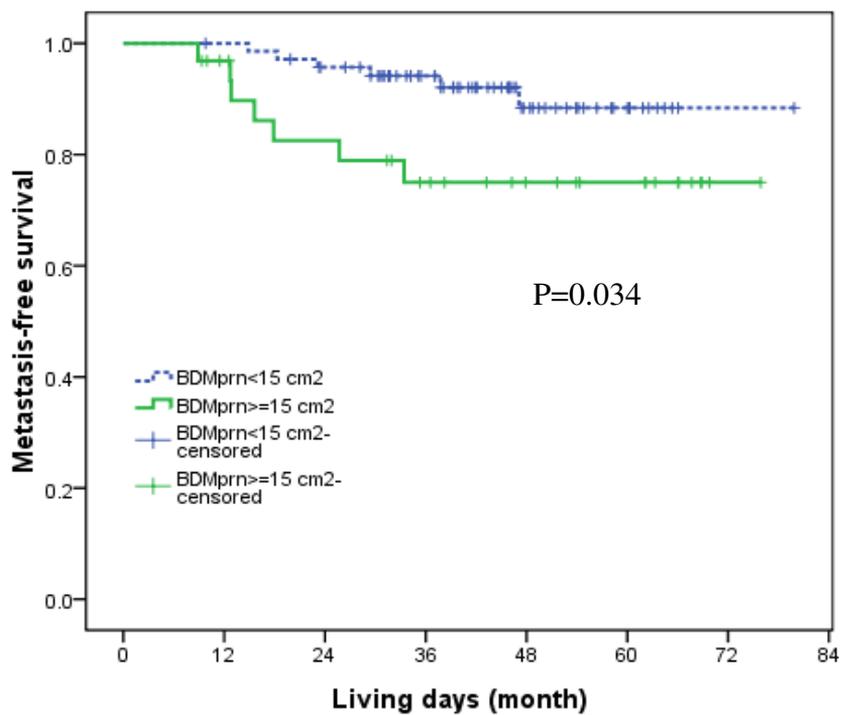


B

Figure 1



A



B

Additional files provided with this submission:

Additional file 1: 2129458047405056 - Chang Add file.docx, 11K
<http://www.ro-journal.com/imedia/2181620354365664/supp1.docx>