

University of Dundee

Nimotuzumab-cisplatin-radiation versus cisplatin-radiation in HPV negative oropharyngeal cancer

Noronha, Vanita; Patil, Vijay Maruti; Joshi, Amit; Mahimkar, Manoj; Patel, Usha; Pandey, Manish Kumar

Published in:
Oncotarget

DOI:
[10.18632/oncotarget.27443](https://doi.org/10.18632/oncotarget.27443)

Publication date:
2020

Licence:
CC BY

Document Version
Publisher's PDF, also known as Version of record

[Link to publication in Discovery Research Portal](#)

Citation for published version (APA):

Noronha, V., Patil, V. M., Joshi, A., Mahimkar, M., Patel, U., Pandey, M. K., Chandrasekharan, A., Dsouza, H., Bhattacharjee, A., Mahajan, A., Sabale, N., Agarwal, J. P., Ghosh-Laskar, S., Budrukkar, A., D'Cruz, A. K., Chaturvedi, P., Pai, P. S., Chaukar, D., Nair, S., ... Prabhash, K. (2020). Nimotuzumab-cisplatin-radiation versus cisplatin-radiation in HPV negative oropharyngeal cancer. *Oncotarget*, 11(4), 399-408. <https://doi.org/10.18632/oncotarget.27443>

General rights

Copyright and moral rights for the publications made accessible in Discovery Research Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Nimotuzumab-cisplatin-radiation versus cisplatin-radiation in HPV negative oropharyngeal cancer

Vanita Noronha^{1,*}, Vijay Maruti Patil^{1,*}, Amit Joshi¹, Manoj Mahimkar², Usha Patel², Manish Kumar Pandey², Arun Chandrasekharan¹, Hollis Dsouza¹, Atanu Bhattacharjee³, Abhishek Mahajan¹, Nilesh Sabale¹, Jai Prakash Agarwal⁴, Sarbani Ghosh-Laskar⁴, Ashwini Budrukkar⁴, Anil K. D'Cruz⁵, Pankaj Chaturvedi⁵, Prathamesh S. Pai⁵, Devendra Chaukar⁵, Sudhir Nair⁵, Shivakumar Thiagarajan⁵, Shripad Banavali¹ and Kumar Prabhash¹

¹Department of Medical Oncology, Tata Memorial Hospital, HBNI, Mumbai, India

²Mahimkar Laboratory, Cancer Research Institute (CRI), Advanced Centre for Treatment Research and Education in Cancer (ACTREC), Tata Memorial Centre (TMC), HBNI, Navi Mumbai, India

³Section of Biostatistics, Department of Epidemiology, Advanced Centre for Treatment, Research and Education in Cancer, Navi Mumbai, India

⁴Department of Radiation Oncology, Tata Memorial Hospital, HBNI, Mumbai, India

⁵Department of Head and Neck Surgery, Tata Memorial Hospital, HBNI, Mumbai, India

*These authors contributed equally to this work

Correspondence to: Kumar Prabhash, **email:** kumarprabhashtmh@gmail.com

Keywords: HPV negative; oropharynx; nimotuzumab; weekly; cisplatin

Received: September 03, 2019

Accepted: December 21, 2019

Published: January 28, 2020

Copyright: Noronha et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License 3.0 (CC BY 3.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

ABSTRACT

Background: Addition of nimotuzumab to weekly cisplatin and radiation improves outcomes in head and neck cancer. HPV negative oropharyngeal cancer has unsatisfactory treatment outcomes and is a candidate for escalation of treatment. We wanted to determine whether the addition of nimotuzumab to cisplatin-radiation could improve outcomes in these poor-risk tumors.

Methods: This was a subgroup analysis of a phase 3 randomized study. In this study, locally advanced head and neck cancer patients undergoing definitive chemoradiation were randomly allocated to weekly cisplatin (30 mg/m² IV)- radiation (66–70 Gy) {CRT arm} or nimotuzumab (200 mg weekly) -weekly cisplatin (30 mg/m²)-radiation (66–70 Gy) {NCRT arm}. The data of HPV negative oropharyngeal cancer was extracted from the database of this study for the analysis. HPV testing was done with p16 immunohistochemistry (IHC) staining and reported according to the CAP criteria. The outcomes assessed were progression-free survival (PFS), disease-free survival (DFS), locoregional control, and overall survival (OS). Interaction test was performed between the study arms and HPV status prior to doing any HPV specific analysis for each of the studied outcomes. Kaplan Meier estimates for 2 year OS with 95%CI was calculated. The hazard ratio was obtained using COX regression analysis.

Results: We had 187 HPV negative oropharyngeal cancers, 91 in the CRT arm and 96 in NCRT arm. The interaction test was significant for PFS ($p = 0.000$), locoregional control ($p = 0.007$) and overall survival ($p = 0.002$) but not for DFS ($p = 0.072$). The 2- year PFS was 31.5% (95%CI 21.5–42) in CRT arm versus 57.2% (95%CI 45.8–67.1) in NCRT arm (HR -0.54; 95%CI 0.36–0.79, $p = 0.002$). The 2-year LRC was 41.4% (95%CI 29.8–52.6) in the CRT arm versus in 60.4% (95%CI 48.7–70.2) in the NCRT arm (HR -0.61; 95%CI 0.4–0.94, $p = 0.024$). The addition of nimotuzumab also lead to an improvement in 2-year OS from 39.0% (95%CI 28.4–49.6) to 57.6% (95%CI 46.3–67.4) (HR-0.63, 95%CI 0.43–0.92, $p = 0.018$).

Conclusions: The addition of nimotuzumab to weekly cisplatin-radiation improves outcomes inclusive of OS in HPV negative oropharyngeal cancers.

INTRODUCTION

Locally advanced oropharyngeal cancers possess a unique challenge. The anatomical location along with multiple physiological functions associated with this structure makes open surgical resection a difficult and moribund procedure [1]. Hence in locally advanced oropharyngeal cancers, concurrent chemoradiation is used primarily for management. The prognosis of these tumors is heavily dependent on the Human Papilloma Virus (HPV) status [2]. Patients with HPV positive disease have a favorable prognosis as opposed to HPV negative disease [2]. The incidence of HPV positive disease is variable across the globe with the incidence in North America in the range of 48–81.4% [3], and 15–22.8% in the Indian subcontinent [4].

Efforts to improve outcomes in locally advanced oropharyngeal cancer have met with limited success [5, 6]. Recently a phase 3 randomized study was reported by us, in locally advanced head and neck cancers, testing the role of concurrent nimotuzumab in addition to weekly cisplatin and definitive radiation. The study met its primary endpoint of progression-free survival [7]. However, a similar study exploring the role of Cetuximab (RTOG 0522) was negative [8]. We had hypothesized that an improvement in outcomes in our study was seen due to the differential patient population in our study when compared to the RTOG 0552 study. We had a younger cohort of patients and a predominantly HPV negative disease [7]. This population, even in RTOG 0522 showed a trend towards improvement with the addition of cetuximab [8]. Another phase 3 study, exploring the role of cetuximab along with carboplatin -5 FU with radiation, reported improvement in outcomes with the addition of cetuximab [9]. The probable reason for this was probably due to the predominant HPV negative disease.

Taking this into consideration, we decided to perform a subgroup analysis of the HPV negative oropharyngeal cancer cohort, to study the absolute improvement in 2-year outcomes with the addition of nimotuzumab. We compared 2 year progression free survival (PFS), disease free survival (DFS), locoregional control (LRC) and overall survival (OS) between both arms.

RESULTS

Patient selection and baseline characteristics

We had 536 patients in the study, out of which 269 had primary in the oropharynx. P16 testing was feasible in 212 patients and 187 patients were p16 negative. These 187 patients are included in the current analysis. There were 91 patients in the cisplatin radiotherapy (CRT) arm and 97 in nimotuzumab-cisplatin radiation (NCRT) arm. The baseline characteristics were balanced between the 2 arms (Table 1).

Outcomes

The interaction test for HPV status (positive & negative) was significant for PFS ($p = 0.000$), LRC ($p = 0.007$) and OS ($p = 0.002$) but not for DFS ($p = 0.072$), suggesting a differential impact of the addition of nimotuzumab with respect to HPV status.

Progression-free survival

At the time of data censoring 103 events had occurred, 60 in CRT arm and 43 in the NCRT arm. The median PFS was 12.9 months (95%CI 8.47–17.3) versus 35.3 months (95%CI 22.10–NA) (P -value = 0.0015) in the CRT and NCRT arm respectively. The 2 year PFS was 31.5% (95%CI 21.5–42) in CRT arm versus 57.2% (95%CI 45.8–67.1) in NCRT arm (Figure 1). The unadjusted hazard ratio was 0.53 (95%CI 0.36–0.79, P -value = 0.002). The multivariate analysis for PFS is shown in Table 2. The adjusted hazard ratio was 0.53 (95%CI 0.36–0.8, P -Value = 0.002). The results of sensitivity analysis performed for PFS using a composite endpoint of progression or death was in line with the above-mentioned analysis. The unadjusted and adjusted hazard ratios were 0.62 (95%CI 0.43–0.88, P -value = 0.008) and 0.62 (95%CI 0.43–0.89, P -value = 0.009), respectively. The site of failure was locoregional in 45 patients (49.5%), locoregional with distant failure in 3 patients (3.3%) and distant failure in 12 patients (13.2%) in CRT arm. The patients with corresponding sites of failure in the NCRT arm were 32 (33.3%), 6 (6.2%) and 5 (5.2%), respectively.

Locoregional control

The improvement in PFS with the addition of nimotuzumab is largely contributed by an improvement in locoregional control. The median time to locoregional failure was 17.3 months (95%CI 12.0–56.3) and 60.3 months (95%CI 22.6–NA) in cisplatin arm and cisplatin-nimotuzumab arm respectively (P -value = 0.023). The 2-year LRC was 41.4% (95%CI 29.8–52.6) in CRT arm versus in 60.4% (95%CI 48.7–70.2) in NCRT arm (Figure 2). The unadjusted hazard ratio was 0.61 (95%CI 0.4–0.94, P -value = 0.024) while the adjusted hazard ratio was 0.63 (95%CI 0.41–0.97, P -value = 0.037). These results were confirmed in the sensitivity analysis. The unadjusted and adjusted hazard ratios were 0.68 (95%CI 0.48–0.99, P -value = 0.0358) and 0.69 (95%CI 0.43–0.89, P -value = 0.0494) in favor of the cisplatin-nimotuzumab arm, with the use of composite endpoint of locoregional failure or death.

Overall survival

The median follow up was 38.5 months (95%CI 36.2–46.3). At the data cutoff, 105 deaths had happened; 57

Table 1: Baseline characteristics

Variable	Cisplatin-radiotherapy arm (n = 91)	Nimotuzumab-cisplatin- radiotherapy arm (n = 96)	P-value
Age, years			
Median age (range)	53 (31–75)	56 (34–70)	
Age > 60 years	23 (25.3)	30 (31.2)	0.272*
Gender			
Male	83 (91.2)	83 (86.5)	0.359
Female	8 (8.8)	13 (13.5)	
ECOG PS			
0	20 (22)	14 (14.6)	0.255
1	71 (78)	82 (85.4)	
Tobacco use			
Yes	85 (93.4)	88 (91.7)	0.783
No	6 (6.6)	8 (9.3)	
Subsite of malignancy			
Base of tongue	52 (57.1)	57 (59.4)	0.959
Tonsil	25 (27.5)	27 (28.1)	
Soft palate	10 (11)	9 (9.4)	
Posterior pharyngeal wall	4 (4.4)	3 (3.1)	
T category [†]			
T1–T2	25 (27.5)	16 (16.7)	0.08
T3–T4	66 (72.5)	80 (83.3)	
N category [†]			
N0–N1	35 (38.5)	35 (36.5)	0.88
N2–N3	56 (61.5)	61 (63.5)	
TNM Stage grouping [†]			
Stage III	14 (15.4)	18 (18.8)	0.794
Stage IVA	74 (81.3)	74 (77.1)	
Stage IVB	3 (3.3)	4 (4.2)	
Histological Grade			
Grade 1–2	68 (74.4)	66 (68.8)	0.418
Grade 3	23 (25.3)	30 (31.2)	

Data are presented as number (%) unless otherwise specified. *P-value provided is for the comparison of age < 60 years versus age > or = 60 years between the 2 arms. [†]The staging was done according to the AJCC-UICC 7th Edition.

in cisplatin arm and 48 in the cisplatin-nimotuzumab arm. The median OS was 19.0 months (95%CI 14.2–23.4) in cisplatin arm while it was 35.9 months (95%CI 22.8–53.7) in the cisplatin-nimotuzumab arm (*P*-value = 0.017). The addition of nimotuzumab also led to an improvement in 2 year OS from 39.0% (95%CI 28.4–49.6) to 57.6% (95%CI

46.3–67.4) (Figure 3). The assumption for proportional hazard was violated (Supplementary Table 1) and hence, COX regression analysis was found to be an unsuitable method for comparison. Alternatively, restricted mean survivals were calculated for both arms and compared. The restricted mean survival estimated in cisplatin and cisplatin-

nimotuzumab arm using data up to 65 months was 37.62 months (95%CI 32.5–42.74) and 29.45 months (95%CI 24.12–34.77), respectively (Figure 4), a difference of 8.17 months (95%CI 0.79–15.56; P -value = 0.03).

Compliance to treatment

The data for treatment compliance and radiation technique are shown in Table 3. There was no difference in these factors between the 2 arms.

Adverse events

The clinical adverse events were captured in 180 patients while laboratory adverse events were captured in 182 patients. The adverse events details between the 2 arms are shown in Supplementary Table 2.

DISCUSSION

The results of the current study clarify the importance of treatment intensification in HPV negative oropharyngeal cancers. Locoregional control, progression-free survival and overall survival were improved with the addition of nimotuzumab to cisplatin and radiation. The absolute improvement in the 2 years LRC and PFS were around 20–25 percent. The corresponding improvement in

OS was 18 percent. These improvements, in accordance with the ESMO magnitude of clinical benefit scale for curative treatment would classify as “A” [10].

As opposed to HPV related oropharyngeal cancer, HPV negative oropharyngeal cancers have worse prognosis. The median PFS, LRC, and OS in the current study are however, lower than reported in literature from the western world [8, 11, 12], [13–15]. The patient population in the current study had a history of tobacco use (90 percent) with the predominance of stage IV disease (80 percent), which probably resulted in lower survival. Both tobacco and stage IV are known bad prognostic factors [2, 16]. Another factor responsible for poorer outcomes would be the use of weekly cisplatin. However, it is unlikely as the cumulative dose of cisplatin received was 200 mg per M2 or more in nearly 3/4th of the patients. Dose intensification above 200 mg/m2 has questionable benefit [17]. These results in the control arm are similar to the results we previously reported [16].

Addition of nimotuzumab as a radiosensitizer to weekly cisplatin and addition of cetuximab to carboplatin 5-FU resulted in an improvement in outcomes over weekly cisplatin and carboplatin-5 FU in locally advanced head and neck cancers [7, 9]. However, a similar study of the addition of cetuximab to 3 weekly cisplatin was associated with negative results [8]. The probable reason for this discrepancy was the nature of the population in these 3

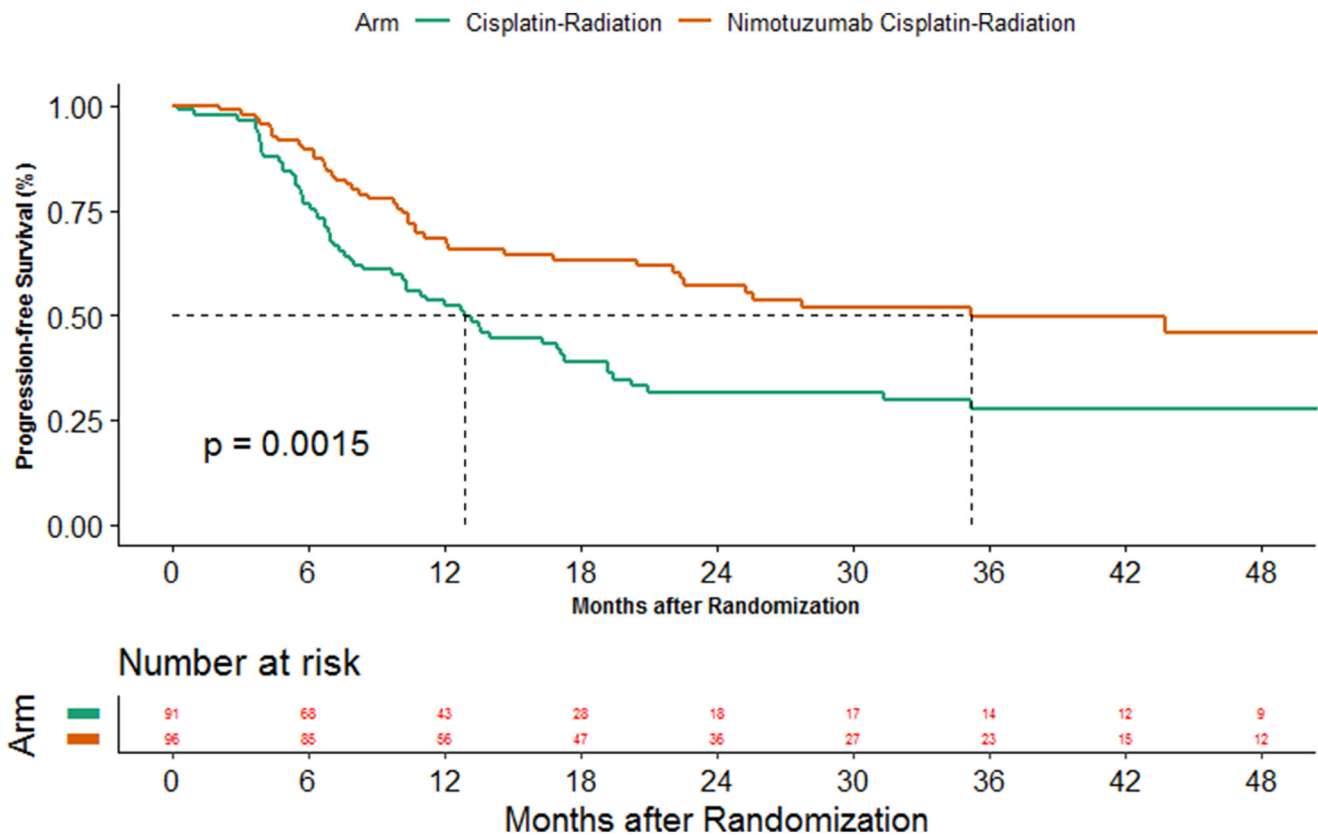


Figure 1: Kaplan Meier estimates of progression-free survival between the 2 arms.

Table 2: Table depicting the result of multivariate cox regression analysis for progression-free survival and locoregional control

Variables	Variable type	Reference	Hazard ratio (HR)	95%CI of HR	P-value
Progression-free analysis					
Arm	Binary	Cisplatin arm	0.5352	0.3588–0.7983	0.00218*
Age	Binary	Below 60 years	0.7138	0.4573–1.1141	0.13776
Stage	Binary	Stage III	1.7770	0.9587–3.2937	0.06786
ECOG PS	Binary	ECOG PS 0	0.8501	0.5337–1.3541	0.49418
Grade	Binary	Grade 1–2	1.2734	0.8227–1.9710	0.27816
Subsite	Binary	Non base of tongue	1.0607	0.7106–1.5835	0.77303
Time to locoregional control					
Arm	Binary	Cisplatin arm	0.6311	0.4094–0.9728	0.0371*
Age	Binary	Below 60 years	0.7268	0.4453–1.1861	0.2016
Stage	Binary	Stage III	2.0823	1.0250–4.2303	0.0425*
ECOG PS	Binary	ECOG PS 0	0.6850	0.4203–1.1165	0.1290
Grade	Binary	Grade 1–2	1.1278	0.6967–1.8256	0.6245
Subsite	Binary	Non base of tongue	0.9464	0.6121–1.4631	0.8042

ECOG PS: Eastern Cooperative Oncology Group Performance Status. *Statistically significant values.

studies. In our study and the french study, the predominant population was HPV negative while, in the RTOG 0552, the predominant population was HPV positive. We feel that dose intensification would work if the population had poor prognostic factors like HPV negative. Hence, this analysis was performed to study whether addition of nimotuzumab would have larger incremental benefits in a HPV negative population. And indeed it leads to larger

incremental benefits as opposed to the whole cohort Supplementary Table 3.

The current study is not without limitations. It is a subgroup analysis. However, an interaction test was performed prior to doing the subgroup analysis, to determine whether nimotuzumab had a differential impact or not. The study had used weekly cisplatin and not 3 weekly cisplatin, and so, the outcomes of the analysis are only applicable for weekly regimen.

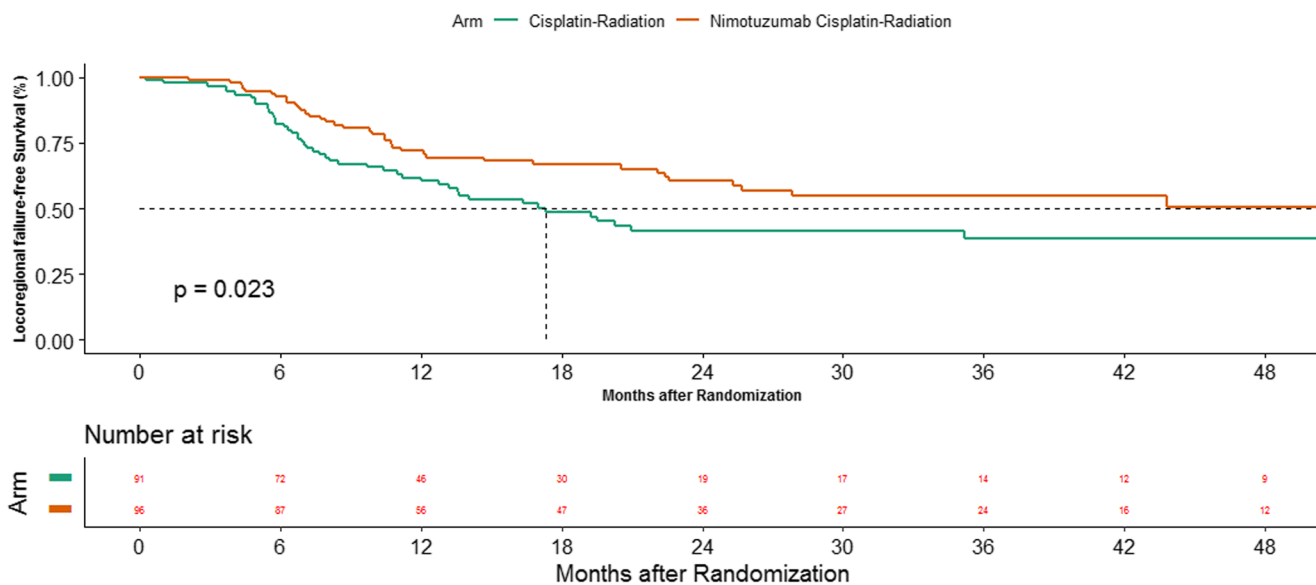


Figure 2: Kaplan Meier estimates of time to locoregional control between the 2 arms.

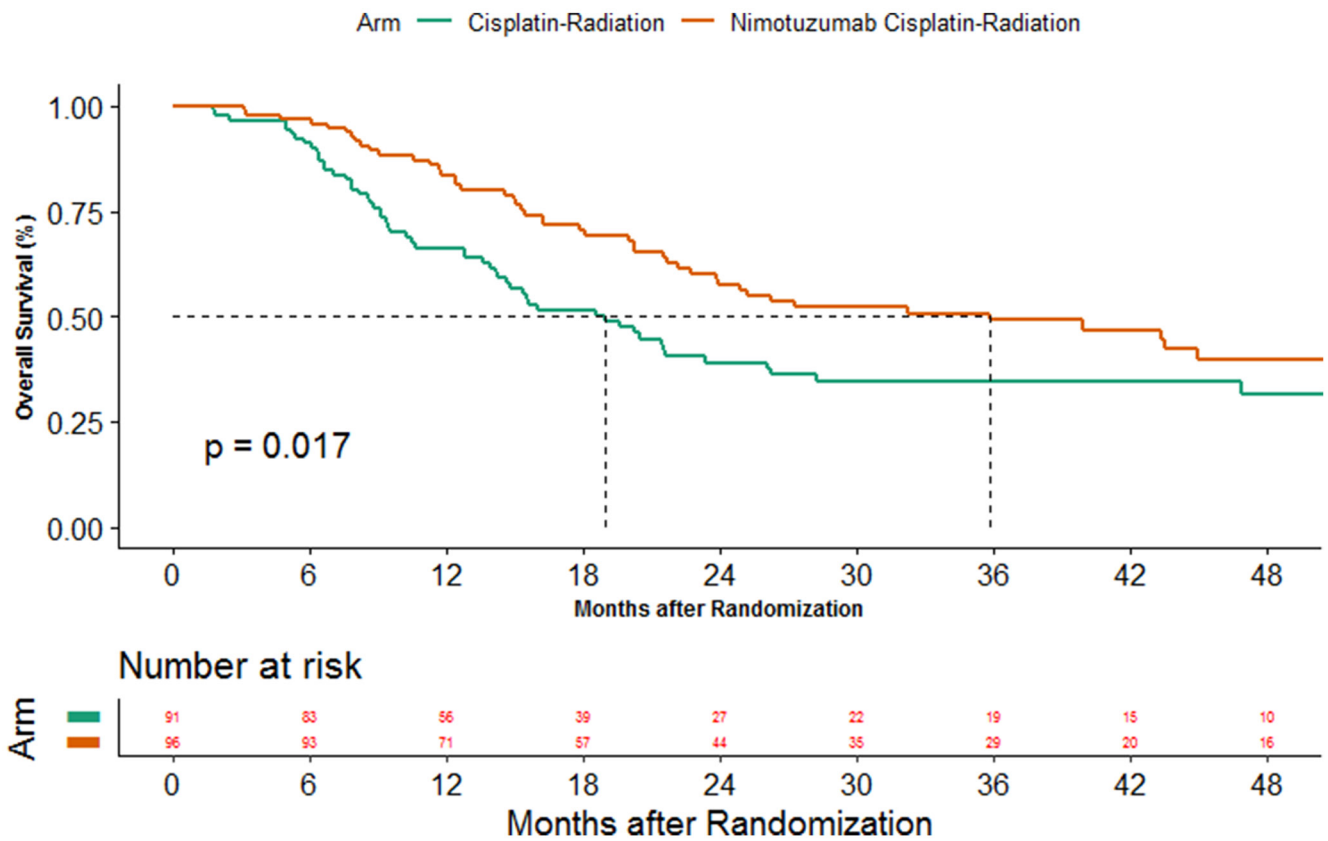


Figure 3: Kaplan Meier estimates overall survival between the 2 arms.

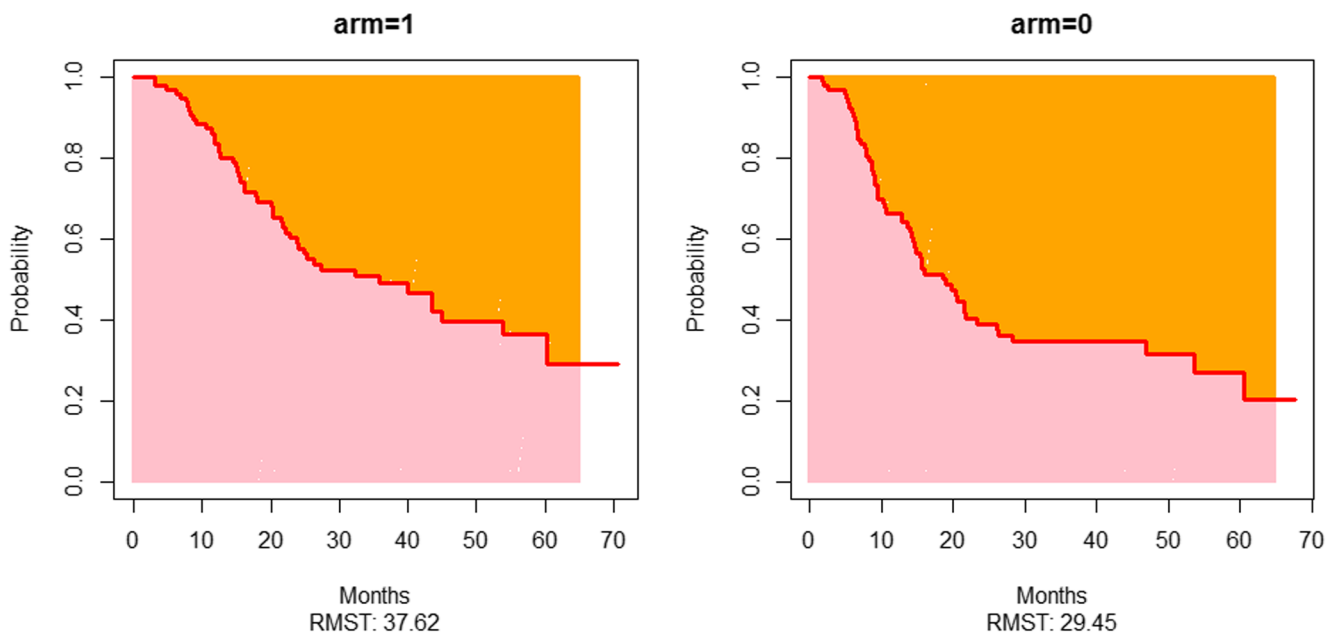


Figure 4: Restricted mean overall survival plots of both arms. arm = 0 represents the plot of the cisplatin radiotherapy arm while arm = 1 represents the plot of the Nimotuzumab-cisplatin radiotherapy arm.

Table 3: Compliance data for radiation and cisplatin

Variable	Cisplatin-radiotherapy arm	Nimotuzumab-cisplatin-radiotherapy arm	P-value
	(n = 91)	(n = 96)	
Radiotherapy compliance			
Radiotherapy dose			
Median dose	70 (IQR 70–70)	70 (IQR 70–70)	–
100%	86 (94.5)	84 (87.5)	0.127
≥95%	86 (94.5)	85 (88.5)	0.193
Radiotherapy technique			
2-Dimensional	83 (91.2)	86 (89.6)	
IMRT	8 (8.8)	9 (9.4)	1.0
Not started	–	1 (1.0)	
Median package time in days	52 (IQR 49–55)	51 (IQR 49–54)	–
Treatment completed within 63 days			
Yes	88 (96.7)	93 (96.9)	1.0
No	3 (3.3)	3 (3.1)	
Gaps			
1 day or more	30 (33)	36 (37.5)	0.543
≥3 days cumulative duration	24 (26.4)	32 (33.3)	0.34
Systemic therapy compliance			
Cisplatin cycles			
Median	7 (IQR 7–7)	7 (IQR 7–7)	
7 or more	74 (81.3)	83 (86.5)	0.426
Cumulative dose 200 mg/m ² or above of cisplatin			
Yes	72 (79.1)	77 (80.2)	0.858
No	19 (20.9)	19 (19.8)	
Cisplatin dose reduction			
Yes	7 (7.7)	11 (11.5)	0.461
No	84 (92.3)	85 (88.5)	
Nimotuzumab			
Median	–	7 (IQR 7–7)	–

MATERIALS AND METHODS

Patient population and design of the study

This was a phase 3 randomized superiority study conducted between 2012–2018. The study protocol was approved by the institutional ethics committee and the

study was conducted in accordance with national (Indian Council of Medical Research) and international guidelines (Good Clinical Practice and Declaration of Helsinki) on human research. The detailed inclusion-exclusion criteria with study protocol are already published [7]. The study enrolled patients with locally advanced head and neck cancer who were planned for curative intent therapy, had

normal organ functions, no uncontrolled comorbidities and ECOG PS 0–2. Patients with primary in salivary gland, nasopharynx or thyroid were excluded. For the current analysis, we selected locally advanced oropharyngeal cancer patients who were considered as p16 negative on immunohistochemistry.

Intervention in both arms

The standard arm received radical radiation to a dose of 66–70 Gy with conventional fractionation over 6–7 weeks. Altered fractionation schedules were allowed in both arms if the Biologically equivalent dose (BED) was around 70Gy10. The chemosensitizer used was weekly cisplatin –30 mg/m². The dose of 30 mg/m² was selected as it was considered as standard at our centre and had been proven effective compared to radical radiation in a randomized study [18]. In the experimental arm, in addition to cisplatin and radiation, nimotuzumab was administered as a flat dose of 200 mg, over 1 hour in 0.9% normal saline without any premedication. Cisplatin in both arms was administered over 1 hour with adequate antiemetic cover and hydration.

HPV testing

HPV testing was done using p16 IHC. The procedure for performing HPV testing has been already published in the literature [19]. Interpretation of p16 was done in accordance with American pathologist consensus guidelines. Both negative and positive controls were performed while performing p16 IHC for each batch for quality assurance.

Study conduct

All patients, post consenting, underwent screening procedures, which were complete blood count, renal function tests, liver function test, viral serology, pure tone audiometry, dental examination, speech – swallowing function testing, nutritional status examination and evaluation by a social worker for compliance. All patients underwent an examination under anesthesia for mucosal disease assessment and a contrast enhanced computed tomography (CECT) neck with upper thorax for staging. In patients who had N2 nodes, the imaging used as positron emission tomography (PET). The radiation was administered either via conventional 2-D technique or 3-D technique. Use of intensity-modulated radiotherapy was permitted as per physician's discretion. The patients were assessed each week during the treatment phase. Post-treatment completion at 8 weeks, patients were assessed in a joint clinic with a PET. Further follow up was according to the study protocol. The patients were followed up until death.

Endpoint definition

The primary endpoint was PFS. It was defined as the time in months from randomization until progression. Progression was defined as per RECIST version 1.1. The secondary endpoints were LRC, DFS and OS. Locoregional control was defined as the time in months from randomization till loco-regional progression. Overall survival was defined as the time in months from randomization till death.

Statistical analysis

Descriptive statistics were performed. Nominal and ordinal data between both arms were compared using Fischer's test. An interaction test was performed between HPV status and outcomes (PFS, DFS, LRC, and OS). Only if the interaction test was positive (*P*-value of 0.05 or below), which suggested that the addition of nimotuzumab had a differential impact on outcome in accordance with HPV status, further analysis was done on HPV negative patients.

The subgroup had 187 patients. Considering a type 1 error of 5% and a type 2 of 20%, this subgroup was powered to rule out a 20% absolute improvement at 2 years in PFS, which was the primary endpoint.

Kaplan Meier method was used for estimation of the probability of PFS, LRC, and OS in each arm. The median estimates with the 95% confidence interval (CI) were reported. The 95%CI was calculated in accordance with Brookmeyer and Crowley method. The unadjusted hazard ratio (HR) and adjusted HR (adjusted for prognostic factors) with its 95%CI interval were calculated using the COX regression analysis with Efron's method of tie handling, with CRT arm being considered as reference. The assumptions of proportional hazard model were checked using Schoenfeld residuals and assumptions were met for PFS and LRC. However, the assumption was violated for OS. Hence mean overall survival between the 2 arms were calculated using the restricted mean survival method. The difference between the restricted means between the 2 arms with its 95%CI were calculated. A *p*-value of 0.05 or below was considered as significant.

CONCLUSIONS

The addition of nimotuzumab to weekly cisplatin-radiation improves outcomes inclusive of OS in HPV negative oropharyngeal cancers and this regimen should be considered as standard if 3 weekly cisplatin 100 mg/m² regimen is not used.

Author contributions

Kumar Prabhash and Vijay Patil had full access to all the data in the study and take responsibility for the

integrity of the data and the accuracy of the data analysis. Data analysis was conducted by Vijay Patil, Atanu Bhattacharjee, and Kumar Prabhash.

CONFLICTS OF INTEREST

The authors declare the following financial interests/ personal relationships which may be considered as potential competing interests. Dr. Noronha reports research grants from Dr. Reddy's Laboratories Inc, Amgen, Sanofi India Ltd., Intas Pharmaceuticals and Astra Zeneca Pharma India Ltd., outside the submitted work. Dr. Prabhash reports grants from Biocon Ltd, grants from Dr. Reddy's Laboratories Inc, grants from Fresenius Kabi India Pvt Ltd, grants from Alkem Laboratories, grants from Natco Pharma Ltd, grants from BDR Pharmaceuticals Intl Pvt Ltd, grants from Roche Holding AG, outside the submitted work. All grants were paid to the institution. None of the other authors have anything to declare that may be considered as potential competing interests.

FUNDING

We thank Biocon Ltd and the Tata Memorial Center Research Administration Council for funding this study. The funding agency had no role in design and conduct of the study, collection, management, analysis, and interpretation of the data, preparation, review or approval of the manuscript, and decision to submit the manuscript for publication.

REFERENCES

1. Parsons JT, Mendenhall WM, Stringer SP, Amdur RJ, Hinerman RW, Villaret DB, Moore-Higgs GJ, Greene BD, Speer TW, Cassisi NJ, Million RR. Squamous cell carcinoma of the oropharynx: surgery, radiation therapy, or both. *Cancer*. 2002; 94:2967–80. <https://doi.org/10.1002/ncr.10567>. [PubMed]
2. Ang KK, Harris J, Wheeler R, Weber R, Rosenthal DI, Nguyen-Tân PF, Westra WH, Chung CH, Jordan RC, Lu C, Kim H, Axelrod R, Silverman CC, et al. Human papillomavirus and survival of patients with oropharyngeal cancer. *N Engl J Med*. 2010; 363:24–35. <https://doi.org/10.1056/NEJMoa0912217>. [PubMed]
3. Stein AP, Saha S, Kraninger JL, Swick AD, Yu M, Lambert PF, Kimple RJ. Prevalence of Human Papillomavirus in Oropharyngeal Cancer: A Systematic Review. *Cancer J*. 2015; 21:138–46. <https://doi.org/10.1097/PPO.000000000000115>. [PubMed]
4. Murthy V, Calcuttawala A, Chadha K, d'Cruz A, Krishnamurthy A, Mallick I, Nair S, Teni T, Pawar S, Talapatra K, Patil A, Bhatt A, Chatterjee S, et al. Human papillomavirus in head and neck cancer in India: current status and consensus recommendations. *South Asian J Cancer*. 2017; 6:93–98. https://doi.org/10.4103/sajc.sajc_96_17. [PubMed]
5. Pignon JP, Baujat B, Bourhis J. [Individual patient data meta-analyses in head and neck carcinoma: what have we learnt?]. [Article in French]. *Cancer Radiother*. 2005; 9:31–36. <https://doi.org/10.1016/j.canrad.2004.11.002>. [PubMed]
6. Nguyen-Tan PF, Zhang Q, Ang KK, Weber RS, Rosenthal DI, Soulieres D, Kim H, Silverman C, Raben A, Galloway TJ, Fortin A, Gore E, Westra WH, et al. Randomized phase III trial to test accelerated versus standard fractionation in combination with concurrent cisplatin for head and neck carcinomas in the Radiation Therapy Oncology Group 0129 trial: long-term report of efficacy and toxicity. *J Clin Oncol*. 2014; 32:3858–66. <https://doi.org/10.1200/JCO.2014.55.3925>. [PubMed]
7. Patil VM, Noronha V, Joshi A, Agarwal J, Ghosh-Laskar S, Budrukkar A, Murthy V, Gupta T, Mahimkar M, Juvekar S, Arya S, Mahajan A, Agarwal A, et al. A randomized phase 3 trial comparing nimotuzumab plus cisplatin chemoradiotherapy versus cisplatin chemoradiotherapy alone in locally advanced head and neck cancer. *Cancer*. 2019; 125:3184–97. <https://doi.org/10.1002/ncr.32179>. [PubMed]
8. Ang KK, Zhang Q, Rosenthal DI, Nguyen-Tan PF, Sherman EJ, Weber RS, Galvin JM, Bonner JA, Harris J, El-Naggar AK, Gillison ML, Jordan RC, Kanski AA, et al. Randomized phase III trial of concurrent accelerated radiation plus cisplatin with or without cetuximab for stage III to IV head and neck carcinoma: RTOG 0522. *J Clin Oncol*. 2014; 32:2940–50. <https://doi.org/10.1200/JCO.2013.53.5633>. [PubMed]
9. Tao Y, Auperin A, Sire C, Martin L, Khoury C, Maingon P, Bardet E, Kaminsky MC, Lapeyre M, Chatellier T, Alfonsi M, Pointreau Y, Jadaud E, et al. Improved Outcome by Adding Concurrent Chemotherapy to Cetuximab and Radiotherapy for Locally Advanced Head and Neck Carcinomas: Results of the GORTEC 2007-01 Phase III Randomized Trial. *J Clin Oncol*. 2018 Jun 7. <https://doi.org/10.1200/JCO.2017.76.2518>. [Epub ahead of print]. [PubMed]
10. Cherny NI, Dafni U, Bogaerts J, Latino NJ, Pentheroudakis G, Douillard JY, Tabernero J, Zielinski C, Piccart MJ, de Vries EG. ESMO-Magnitude of Clinical Benefit Scale version 1.1. *Ann Oncol*. 2017; 28:2340–66. <https://doi.org/10.1093/annonc/mdx310>. [PubMed]
11. Cohen EE, Karrison TG, Kocherginsky M, Mueller J, Egan R, Huang CH, Brockstein BE, Agulnik MB, Mittal BB, Yunus F, Samant S, Raez LE, Mehra R, et al. Phase III randomized trial of induction chemotherapy in patients with N2 or N3 locally advanced head and neck cancer. *J Clin Oncol*. 2014; 32:2735–43. <https://doi.org/10.1200/JCO.2013.54.6309>. [PubMed]
12. Haddad R, O'Neill A, Rabinowits G, Tishler R, Khuri F, Adkins D, Clark J, Sarlis N, Lorch J, Beitler JJ, Limaye S, Riley S, Posner M. Induction chemotherapy

- followed by concurrent chemoradiotherapy (sequential chemoradiotherapy) versus concurrent chemoradiotherapy alone in locally advanced head and neck cancer (PARADIGM): a randomised phase 3 trial. *Lancet Oncol.* 2013; 14:257–64. [https://doi.org/10.1016/S1470-2045\(13\)70011-1](https://doi.org/10.1016/S1470-2045(13)70011-1). [PubMed]
13. Ghi MG, Paccagnella A, Ferrari D, Foa P, Alterio D, Codecà C, Nolè F, Verri E, Orecchia R, Morelli F, Parisi S, Mastromauro C, Mione CA, et al, and GSTTC (Gruppo di Studio Tumori della Testa e del Collo) Italian Study Group. Induction TPF followed by concomitant treatment versus concomitant treatment alone in locally advanced head and neck cancer. A phase II-III trial. *Ann Oncol.* 2017; 28:2206–12. <https://doi.org/10.1093/annonc/mdx299>. [PubMed]
 14. Bonner JA, Harari PM, Giralt J, Cohen RB, Jones CU, Sur RK, Raben D, Baselga J, Spencer SA, Zhu J, Youssoufian H, Rowinsky EK, Ang KK. Radiotherapy plus cetuximab for locoregionally advanced head and neck cancer: 5-year survival data from a phase 3 randomised trial, and relation between cetuximab-induced rash and survival. *Lancet Oncol.* 2010; 11:21–28. [https://doi.org/10.1016/S1470-2045\(09\)70311-0](https://doi.org/10.1016/S1470-2045(09)70311-0). [PubMed]
 15. Rischin D, Peters LJ, O’Sullivan B, Giralt J, Fisher R, Yuen K, Trotti A, Bernier J, Bourhis J, Ringash J, Henke M, Kenny L. Tirapazamine, cisplatin, and radiation versus cisplatin and radiation for advanced squamous cell carcinoma of the head and neck (TROG 02.02, HeadSTART): a phase III trial of the Trans-Tasman Radiation Oncology Group. *J Clin Oncol.* 2010; 28:2989–95. <https://doi.org/10.1200/JCO.2009.27.4449>. [PubMed]
 16. Agarwal JP, Mallick I, Bhutani R, Ghosh-Laskar S, Gupta T, Budrukkar A, Murthy V, Sengar M, Dinshaw KA. Prognostic factors in oropharyngeal cancer—analysis of 627 cases receiving definitive radiotherapy. *Acta Oncol.* 2009; 48:1026–33. <https://doi.org/10.1080/02841860902845839>. [PubMed]
 17. Ang KK. Concurrent radiation chemotherapy for locally advanced head and neck carcinoma: are we addressing burning subjects? *J Clin Oncol.* 2004; 22:4657–59. <https://doi.org/10.1200/JCO.2004.07.962>. [PubMed]
 18. Ghosh-Laskar S, Kalyani N, Gupta T, Budrukkar A, Murthy V, Sengar M, Chaukar D, Pai P, Chaturvedi P, D’Cruz A, Agarwal J. Conventional radiotherapy versus concurrent chemoradiotherapy versus accelerated radiotherapy in locoregionally advanced carcinoma of head and neck: results of a prospective randomized trial. *Head Neck.* 2016; 38:202–07. <https://doi.org/10.1002/hed.23865>. [PubMed]
 19. Bhosale PG, Pandey M, Desai RS, Patil A, Kane S, Prabhash K, Mahimkar MB. Low prevalence of transcriptionally active human papilloma virus in Indian patients with HNSCC and leukoplakia. *Oral Surg Oral Med Oral Pathol Oral Radiol.* 2016; 122:609–618.e7. <https://doi.org/10.1016/j.oooo.2016.06.006>. [PubMed]