



# Head and Neck Cancer Treatment Summary

Patient Name: \_\_\_\_\_

DOB: \_\_\_\_\_

MRN: \_\_\_\_\_

Telephone #: \_\_\_\_\_

Email: \_\_\_\_\_

## Care Team

| Provider | Name | Telephone Number |
|----------|------|------------------|
|----------|------|------------------|

Nurse Navigator  
 Head and Neck Surgeon  
 Radiation Oncologist  
 Medical Oncologist  
 Reconstructive Surgeon  
 Primary Care Physician  
 Dentist

Pain Management  
 Speech Pathologist  
 Social Worker  
 Nutrition Support  
 Other:  
 Other:

## Tumor Characteristics

Date of Pathologic Diagnosis: \_\_\_\_\_ Date of Completion of Definitive Therapy: \_\_\_\_\_

Side:  Left  Right  Bilateral  N/A TNM Stage: T \_\_\_\_\_ N \_\_\_\_\_ M \_\_\_\_\_

Subsite:  Lip  Buccal Mucosa  Floor of Mouth  Alveolar Ridge  Oral tongue  Retromolar Trigone  
 Hard Palate  Nasal Cavity  Paranasal Sinus (Specify \_\_\_\_\_)  Nasopharynx (EBV +/-)  
 Soft Palate  Tonsil  Base of Tongue  Cervical Esophagus  Supraglottic Larynx  Glottic Larynx  
 Subglottic Larynx  Pyriform Sinus  Posterior Pharyngeal Wall  Parotid Gland  
 Submandibular Gland  Facial Skin  Neck Skin  Scalp Skin  Unknown Primary  
 Other: \_\_\_\_\_

Pathology:  Squamous Cell Carcinoma  Other \_\_\_\_\_ HPV/p16 Status:  +  -  N/A

Adverse Pathologic Features:  Positive Margins  Close Margins\*  Perineural Invasion  
 Lymphovascular Invasion  Extracapsular Nodal Extension  
 Skullbase Invasion  Soft tissue invasion (List: \_\_\_\_\_)  
 Other \_\_\_\_\_

\*The designation 'Close margin' is ill-defined and may be used for cancer-free margins from 2-5 mm

## Treatment Details

### Surgery

Surgery Performed:  Yes  No Procedure Date(s): \_\_\_\_\_

Treating Institution: \_\_\_\_\_

Procedure(s) Performed: \_\_\_\_\_

### Radiation

Radiation Therapy Performed:  Yes  No Start Date: \_\_\_\_\_ End Date: \_\_\_\_\_

Treating Institution: \_\_\_\_\_

| Field | Dose | Notes: |
|-------|------|--------|
|       |      |        |
|       |      |        |
|       |      |        |

Chemotherapy Administered:  Yes  No

| Drug Name | Route | Dose | Schedule | Dose reduction   | # Cycles |
|-----------|-------|------|----------|--|----------|
|           |       |      |          | <input type="checkbox"/> Yes _____ % <input type="checkbox"/> No |          |
|           |       |      |          | <input type="checkbox"/> Yes _____ % <input type="checkbox"/> No |          |

### Other

Clinical Trial Participant:  Yes  No Sponsor Name: \_\_\_\_\_ Identifier/NCT #: \_\_\_\_\_

Brief Description: \_\_\_\_\_

Other Treatments Received: \_\_\_\_\_