Perineural Invasion of Skin Cancers in the Head and Neck: An Uncommon Phenomenon Revisited

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Abstract

Objective: The purpose of this article is to describe the epidemiology, imaging findings, pathogenesis, and clinical impact of perineural invasion of skin cancers in the head and neck.

Conclusion: Perineural invasion in head and neck skin cancer can be microscopic disease discovered on pathology or gross perineural spread that can be predicted on imaging often accompanied with clinical symptoms. Physicians and radiologists should have high index of suspicion when evaluating with patient with skin cancer that is in close proximity to a cranial nerve. The advancement of imaging techniques has improved pre-operative detection of perineural invasion of skin cancer.

Keywords: Perineural invasion; Head and neck skin cancer; MRI

Introduction

Perineural Invasion (PNI) of tumor cells was first discovered by Cruveilhier in 1835, when he reported mammary carcinoma invading the facial nerve. In 1862, Neumann first described lower face skin cancer invading the mental nerve.

PNI describes a microscopic finding of tumor infiltration along a nerve, and is distinguished from Perineural Spread (PNS), which describes the presence of gross tumor growth along a nerve distinct from the main tumor mass on imaging. PNI is defined as cancer cell invasion in, around, and through nerves where tumor cells are seen within any of the layers (epineurium, perineurium, endoneurium) of the nerve sheath [1]. When tumor cells are not seen in the layers of nerve sheath, it can be difficult to differentiate it from tumor abutting on nerve.

PNI of tumor is a well-recognized mechanism of tumor dissemination. PNI from head and neck cancer is well-known disease entity, ranging from 27-82%, depending on histology and type of cancer [2]. Within the head and neck, adenoid cystic cancer has a high propensity for PNI, and squamous cell carcinoma can be often associated with PNI. Presence of PNI has a major negative impact on management and prognosis of patients with Hand N cancer. PNI from cutaneous malignancy is under-recognized and frequently overlooked clinically and radio graphically. Most common type of skin cancers with PNI are Squamous Cell Cancer (SCC), Basal Cell Cancer (BCC) and Neurotrophic Malignant Melanoma (NMM). The reported incidences of PNI in non-melanoma skin cancers are 0.18% to 10% in BCC and 2.5% to 14% in SCC [3,4]. The concept of “skip lesions” in PNI is controversial. One set of authors believe in the presence of so called skip lesions in PNI, although none of them provide a convincing evidence to prove the presence of this entity [5-8]. The other set of authors believe that the concept of skip lesions in PNI does not exist and is a misinterpretation which is being propagated in the medical literature by blind quoting of existing false information in the medical literature [9-11]. PNI should be differentiated from neural clinical manifestations that are due to gross extrinsic compression by a tumor on the nerve or direct tumor extension along the skull base foramina or fissures through different anatomical planes [10]. The cranial nerve most vulnerable for PNI from skin cancer is the trigeminal nerve due to its rich cutaneous innervation in the region of most UV exposed regions of the head and neck [12]. The facial nerve is also commonly involved with PNI, especially in cases of SCC that are metastatic to the parotid gland. Less commonly, greater auricular nerve from cervical plexus and nerves of extra-ocular movement (III, IV, and VI) can be involved.

Relevant Cranial Nerve Anatomy

Trigeminal nerve (V cranial nerve)

The trigeminal nerve is a mixed nerve with both motor and sensory components and is considered to be the largest of all cranial nerves. There are four cranial nerve V nuclei, mesencephalic nucleus (proprioception), main sensory (tactile sensation), main motor nucleus (motor innervation to the muscles of mastication) and the spinal nucleus (primitive sensations like pain and temperature), which span across the inferior midbrain to the upper cervical cord to the level of C2. The entry point of the cisternal segment of the nerve in the lateral aspect of pons is called ‘root entry zone’ (REZ). The nerve then relays in the gasserian ganglion located in the Meckel cave, which then divides in to V1 (Ophthalmic), V2 (Maxillary) and V3 (Mandibular) divisions. V1 and V2 divisions are pure sensory nerves and transit through the lateral wall of cavernous sinus and exit the cranial cavity through the superior orbital fissure (in to orbit) and foramen rotundum (in to pterygopalatine fossa), respectively. V1 and V2 divisions of trigeminal nerve innervate upper and mid 1/3rd of the face, respectively. The V3 division is a mixed nerve with motor innervation to the muscles of mastication and sensory to the lower 1/3rd of the face. It exits the cranial cavity through the foramen ovale in to the masticator space (Figure 1a).

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Facial nerve (VII cranial nerve)

The facial nerve is a mixed cranial nerve with motor, parasympathetic and taste sensory components in it. It has three nuclei (motor nucleus, superior salivatory and solitarius tract nucleus) located in the inferior pons antero-lateral to the abducens nucleus. The motor fibers arch around the ipsilateral abducens nucleus in the brainstem forming a bulge in the floor of IV ventricle called facial colliculus. The nerve then exits in to the cerebellopontine cistern at the pontomedullary junction and travels along the anterior and superior part of Internal Auditory Canal (IAC) to enter the petrous bone. The petrous portion of the nerve is divided into various segments such as labyrinthine segment, geniculate ganglion, tympanic segment, and mastoid segments. The nerve then exits the skull base through stylomastoid foramen in to the parotid gland, which divides in to five major branches including temporal, zygomatic, buccal, marginal mandibular, and cervical (Figure 1b).

Pathogenesis

The pathogenesis of PNI is unclear. Recently Liebig et al. [14] described PNI as a process resulting due to an active and reciprocal interaction between the nerve endings and the tumor cells. Tumor microenvironment plays a crucial role in the PNI of tumor cells (Figure 2). Few of the important molecular factors involved in this cross talk between the nerve endings and tumor cells include neurotropic factors.

Figure 1a: Sketch diagram of trigeminal nerve showing different segments and their branches.

Figure 1b: Sketch diagram of facial nerve showing different segments and their branches.

Figure 2: Sketch diagram showing interaction between the nerve fibers and tumor cells in the pathogenesis of perineural invasion. Molecules involved in this interaction include the chemokine Fractalkine/Neurotactin (CX3CL1) expressed by neurons and its receptor CX3CR1 on tumor cells and membrane-bound glycoproteins (SIGLEC-4a) binding to tumor mucins (MUC-1). Neurotropins secreted by neural and tumor cells sustain growth and survival of both intratumoral nerves and cancer cells. Colony stimulating factors (G and GM-CSF) secreted by tumor cells sensitize neural cells expressing their receptors (CSFR), affecting pain perception.
like Nerve Growth Factor (NGF), Brain Derived Nerve Growth Factor (BDNF), neurotropins (NT 3, 4 and 5), growth factors and axonal guidance molecules [18,19]. Neurotropins up regulate the tumor cells as well as the intratumoral nerves. Hematopoietic colony stimulating factors (G-CSF and GM-CSF) play a role in tumor-nerve interaction and most importantly tumor induced pain [20]. Myelin-associated glycoprotein (MAC, Siglec-4a) present on the nerve endings and laminin-5 on the tumor cells interact and facilitate PNI of tumor cells [21].

**Diagnosis**

PNI of head and neck skin cancer is often a misdiagnosed entity. The tumor cells in the initial stage invade the small peripheral cutaneous nerves which eventually spread centrally along the larger named nerves and can reach the brainstem if untreated. Depending on the presence or absence of clinical symptoms at the time of diagnosis, it can be classified in to two categories ‘incidental PNI’ and ‘clinical PNI’. Factors predicting the PNI of skin cancer include type of tumor, size of the primary tumor, location, recurrent versus primary untreated tumor and tumor cell differentiation. The incidence of incidental PNI is more common in basal cell cancer as this represents the majority of non-melanoma skin cancer, whereas the clinical PNI is more common in patients with squamous cell cancer [3,22]. Tumors greater the 2 cm located in head and neck region particularly in the mid face and lip region due to rich cutaneous innervation are more vulnerable for PNI [6,23]. Recurrent skin cancers with prior resection and/or radiation therapy are more commonly associated with PNI than the primary untreated skin cancer and are usually of a higher histological grade (histologic grade 3 or 4) [24]. There is also increased incidence of PNI in male patients. Presence of PNI has been linked to increased incidence of regional lymph node, distant metastases and poorer prognosis [25-29].

PNI is definitively diagnosed upon histologic examination of a resected skin cancer or biopsy specimen. Most commonly, PNI is “incidental” or microscopic (mpPNI); a pathological diagnosis made on specimen without the presence of preoperative symptoms suggestive of PNI. But PNI can clinically be suspected (clinical PNI, cPNI) when patients with cutaneous carcinomas manifest neural symptoms such as paresthesia, hypesthesia, pain in the distribution of a trigeminal nerve branch, or facial weakness. Early symptoms can be very subtle and often described as crawling of ants underneath the skin (formication) [30,31]. Early symptoms may go un-noticed, unless the clinician has a very high index of suspicion for PNI. If neglected, symptoms progress to pain, numbness and/or motor deficits along the distribution of the affected cranial nerve. Clinical symptoms may be erroneously ascribed to Bell palsy or trigeminal neuralgia which may further delay the diagnosis by 6 months to 2 years, but can be demonstrated radiographically as described below [12].

Histologically, tumors cells are usually seen invading the nerve or may even extend along the length of the nerve. The tumor cells can spread retrograde toward central brain or brainstem or antegrade toward the distal nerve [32,33]. Occasionally, biopsy of the suspected cranial nerve may be necessary to confirm the diagnosis when clinically suspected preoperatively [34]. Such a biopsy is typically done at the time of resection of the cancer in attempts to confirm a clear margin of resection. Rarely, presence of concentric layers of fibrosis (peritumoral fibrosis) around the tumor cells or surrounded by the tumor cells can mimic PNI and make the diagnosis difficult [35].

**Imaging**

In patients with clinical symptoms suggestive of trigeminal or facial nerve involvement, imaging can be useful to confirm clinical suspicion and guide treatment planning. If neural involvement is confirmed or highly suggestive upon imaging, the extent of surgical resection to obtain a clear surgical margin, including sacrifice of portions of the facial and or trigeminal nerve, may be determined. If imaging demonstrates intracranial PNI or PNS, a lesion may be deemed to be surgically unresectable, and a primary radiation therapy treatment approach that encompasses the path of neural involvement may be considered. In addition, preoperative imaging can guide the extent radiotherapy required in the adjuvant setting. In patients with mpPNI, imaging should be obtained if large nerve involvement is noted in the surgical specimen.

Contrast enhanced MRI is the best imaging study for the detection of PNS and also to assess the extent of disease spread along the cranial nerves. Subtle enhancement of nerve is difficult to diagnose on contrast enhanced CT until it creates a mass lesion. Hi-resolution focused gadolinium enhanced MRI or MR neurography are considered to be the most sensitive imaging for detection of PNI [31]. The MR imaging features that suggest PNI include obliteration of fat plane surrounding the cranial nerve, enhancement with or without enlargement of the nerve, mass in the cavernous sinus or Meckel cave and sometimes changes related to denervation of a group of muscles innervated by the affected cranial nerve suggest indirect signs of PNI (Figures 3-7).

Although the accuracy of MR neurography is described as high as 83% in defining the extent of cranial nerve involvement [36], false negative MRI has been reported ranging from 22-47% of patients with clinical PNI in the early stages [31]. CT can detect foraminarial widening and bony erosion in advanced cases [36-38] (Figure 8).

It is important to thoroughly evaluate cranial nerves that are suspected for invasion on MR imaging. Non contrast T1 weighted images often reveal loss of fat plane in pterygopalatine fossa or mandibular canal. Obliteration of Meckel’s cave or infra orbital foramen also indicates presence of PNS. Asymmetric enhancement of nerve is better appreciated on post contrast MR images with fat suppression. Finally, indirect evidence of PNS can be seen as denervation changes of muscles that are supplied by the suspected cranial nerve (please find examples). In 2001, Williams et al. [39], described the zonal system (Table 1) to

![Figure 3a: 59 year old male patient with prior history of right forehead squamous cell cancer with new onset right forehead numbness and tingling sensation. A: STIR coronal image of orbit demonstrates thickening of the right V1 (ophthalmic) nerve (arrow) and also note the superior ophthalmic vein (SOV) flow void (arrow head).](image-url)
Figure 3b and c: Coronal and axial post Gad fat-suppressed T1WI demonstrates thickening and enhancement of the V1 nerve (arrow) suggesting perineural invasion. Enhancing SOV (arrow head) inferior to the superior rectus muscle.

Figure 4: 72 year old male patient with right cheek squamous cell cancer. A & B: Axial and coronal post Gad fat-suppressed T1WI demonstrates thickening and enhancement of the right V2 (Maxillary) nerve (block arrow) at the foramen rotundum extending in to the cavernous sinus indicating perineural invasion.

Figure 5: 45 year old female patient with history of right lower face metastatic melanoma. A: Coronal fat-suppressed post Gad T1WI demonstrates enlargement of right foramen ovale (curved arrow) with thickening and enhancement of the V3 (mandibular) nerve and right cavernous sinus mass (block arrow) due to perineural invasion. Also note diffuse enhancement of the masticator muscles (arrows) due to acute denervation. B: Axial STIR image showing increased T2 signal in the right masticator muscle due to acute denervation.

Figure 6: 65 year old male patient with past history of cutaneous squamous cell cancer in the right parotid region with intra-parotid recurrence. A: Axial T1WI demonstrate iso-intense mass in the right parotid region with loss of infiltration of fat in the right stylomastoid foramen (yellow arrow). Note the normal fat in left stylomastoid foramen (white arrow). B: Post Gad fat suppressed axial T1WI demonstrate enhancement in the right stylomastoid foramen suggesting perineural invasion along the VII (Facial) nerve and note the lack of enhancement in the normal left stylomastoid foramen (curved arrow). Infiltrating mass in the right masticator space (arrow head) which extended along the V3 nerve through the auriculotemporal nerve.
define the extent of the PNI along the cranial nerve to help guide the treatment options and correlate with the prognosis.

**Treatment and Prognosis**

PNI is most commonly diagnosed incidentally (mPNI), and thus almost all patients have had surgery for the primary tumor resection at the time of diagnosis. Patients with negative surgical margins and focal PNI in small cutaneous nerves (less than 0.1 mm) may be adequately followed clinically without any additional resection, radiation therapy or additional imaging to help determine if more proximal nerve involvement is present. But if patients have other risk factors such as large tumor size, chronic immunosuppression, lymphovascular invasion, large nerve (greater than 0.1 mm) involvement, MRI for to search for additional nerve involvement should be considered as should adjuvant radiation therapy to the primary site and draining nodal basins. Since patients with PNI are at increased risk of nodal metastases up to 15% to 20% [31,40]. For patients with questionable positive PNI margins within the resected specimen, re-excision or adjuvant radiation therapy is recommended, and adjuvant radiation therapy should definitely be recommended in the setting of positive margins, gross residual disease and/or extensive PNI. Hyperfractionation therapy is often used to reduce the risk of radiation induced complications [41].

There is no proven benefit of adjuvant chemotherapy in clinical PNI for skin cancer [42].

In patients with cPNI, treatment depends on the resectability of the primary tumor which is determined with the aid of MR imaging. If the tumor is technically resectable then treatment is usually surgery followed by post-operative radiation. In 2012, Panizza B et al. [9] evaluated 21 patients with head and neck cutaneous SCC and concluded that surgical resection with negative margins in patients with disease extending up to Gasserian or geniculate ganglion provides best chance of cure.

Compared to patients without PNI, PNI is associated with a higher rate of recurrence and 5-year disease specific death. Jambusaria et al. [43] reported a 5-year disease specific survival of 84% in patients with PNI versus 96% in those without. Local site relapse is most common, and most relapses occur within two to four years after initial resection [43]. Clinical PNI patients have an increased rate of relapse compared to those with mPNI. Garcia-Serra et al, reported local control, cause specific survival and overall survival of 87%, 65% and 50% respectively over a 5-year follow up on 59 patients with mPNI compared to 55%, 59% and 55% on 76 patients with cPNI treated with surgery and postoperative radiotherapy[40]. Imaging positive cPNI patients have highest rates of local recurrence, 43% - 75% versus 24% and lower rates of disease-specific survival, 56% - 61% versus 100% compared to image negative patients [12].

**Conclusion**

The radiologist and the clinician should maintain a high index of
susception of perineural involvement when evaluating skin cancers, particularly when they are recurrent and occur in close proximity with one or more cranial nerves in the head and neck region. Patients with skin cancers that exhibit PNI are more likely to exhibit local recurrence and have decreased disease-specific survival after treatment.

References


