Esophageal squamous cell carcinoma (ESCC) is one of the most aggressive squamous cell carcinomas and is highly prevalent in Asia. Alcohol and its metabolite, acetaldehyde, are considered definite carcinogens for the esophagus. Polymorphisms in the aldehyde dehydrogenase 2 gene, which encodes an enzyme that eliminates acetaldehyde, have been associated with esophageal carcinogenesis. Studies of the mutagenic and carcinogenic effects of acetaldehyde support this observation. Several recent large-scale comprehensive analyses of the genomic alterations in ESCC have shown a high frequency of mutations in genes such as TP53 and others that regulate the cell cycle or cell differentiation. Moreover, whole genome and whole exome sequencing studies have frequently detected somatic mutations, such as G:C→A:T transitions or G:C→C:G transversions, in ESCC tissues. Genomic instability, caused by abnormalities in the Fanconi anemia DNA repair pathway, is also considered a pathogenic mechanism of ESCC. Advances in diagnostic techniques such as magnifying endoscopy with narrow band imaging or positron emission tomography have increased the accuracy of diagnosis of ESCC. Updated guidelines from the National Comprehensive Cancer Network standardize the practice for the diagnosis and treatment of esophageal cancer. Patients with ESCC are treated endoscopically or with surgery, chemotherapy, or radiotherapy, based on tumor stage. Minimally invasive treatments help improve the quality of life of patients who undergo such treatments. We review recent developments in the diagnosis and treatment of ESCC and advances gained from basic and clinical research.

**Epidemiology**

The incidence of esophageal cancer is 3-fold higher in men than women, and approximately 80% of cases occur in developing countries. The incidence of esophageal cancer varies greatly with location. There is a high prevalence of ESCC in east Asia, eastern and southern Africa, and southern Europe. In contrast, the incidence of ESCC is low in North America and other parts of Europe. These variations indicate that ethnic and genetic factors and lifestyle all have roles in the development of ESCC.

Alcohol consumption and tobacco use are established risk factors for ESCC and have synergistic effects on risk. Acetaldehyde associated with alcohol intake was added as a definite carcinogen (a group 1 carcinogen) for the esophagus by the International Agency for Research on Cancer. Purported risk factors for ESCC include low levels of consumption of fruits and vegetables; deficiency of selenium, and the 6th leading cause of cancer-related mortality in the world with an estimated 456,000 new cases per year worldwide. Esophageal cancer comprises 2 main histological subtypes: esophageal squamous cell carcinoma (ESCC) and esophageal adenocarcinoma.

**Keywords:** Acetaldehyde; Aldehyde Dehydrogenase 2; Genetic Polymorphism; Field Cancerization; Squamous Differentiation.
Alcohol, Acetaldehyde, and Carcinogenesis

Ingested alcohol is absorbed from the upper gastrointestinal tract and transported to the liver, where it is metabolized to acetaldehyde (intrinsic acetaldehyde) by alcohol dehydrogenase 1B (ADH1B). Acetaldehyde is subsequently detoxified to acetic acid by aldehyde dehydrogenase 2 (ALDH2) (Figure 1). ALDH2 activity is reduced by the polymorphism Glu504Lys,16 which is prevalent in Mongoloid but not in Caucasoid or Negroid populations17; it increases blood, salivary, and breath levels of acetaldehyde after alcohol intake.18,19 Heavy ingestion of alcohol increases the risk of ESCC in people with the ALDH2 Glu504Lys polymorphism.20 This finding could account for the higher incidence of ESCC in Asian versus Western countries.

Acetaldehyde is an organic compound found in ripe fruits, bread, coffee, cheese, and yogurt,21 alcoholic beverages,22 and tobacco smoke.23 Alcoholic beverages such as Calvados and other spirits contain particularly high amounts of acetaldehyde, and frequent consumption of these beverages is associated with an increased risk of ESCC.23 Acetaldehyde can also be generated in the human oral cavity by microorganisms such as yeasts and bacteria.24 Acetaldehyde from alcoholic beverages, foods, and/or oral microflora could therefore provide a potential source of carcinogens that predispose people to ESCC (Figure 1).

Numerous in vitro and in vivo studies support the mutagenic and carcinogenic effects of acetaldehyde.25 Acetaldehyde causes single- and double-strand DNA breaks,26 point mutations,27 sister chromatid exchanges, and gross chromosomal aberrations.28,29 Acetaldehyde also binds to proteins to cause structural and functional alterations.27 Many of the altered enzymes are involved in DNA repair, DNA methylation, and the antioxidative defense system.30 Acetaldehyde causes nasal carcinoma and respiratory squamous cell carcinoma (SCC).31,32 Studies of inhalation in rats and hamsters show that acetaldehyde causes nasal carcinoma and respiratory squamous cell carcinoma (SCC).33,34

Acetaldehyde reacts with DNA to form DNA adducts.35 A single molecule of acetaldehyde react with deoxyguanosine (dG) to generate \( N^2 \)-ethylidene-2′-dG, which can be stabilized by reduction to the product \( N^2 \)-ethyl-2′-dG (Figure 2).36 Because \( N^2 \)-ethylidene-2′-dG is the direct and most abundant DNA adduct derived from acetaldehyde, it is a specific biomarker for identifying acetaldehyde-derived DNA damage.37 \( N^2 \)-ethyl-2′-dG inhibits trans-lesion DNA synthesis, which leads to replication errors and/or frameshift deletion mutations.38 Another class of DNA adducts, \( N^2 \)-propano-2′-dG, which is derived from 2 molecules of acetaldehyde (Figure 2), has also been shown to be mutagenic.39,40 However, the mechanistic role of DNA adducts in acetaldehyde-related ESCC carcinogenesis is unclear.16

Development of ESCC

ESCC develops via a multistep process that begins with a normal squamous epithelium and progresses to low-grade intraepithelial neoplasia (LGIN), high-grade intraepithelial neoplasia (HGIN), and ultimately to invasive carcinoma (Figure 3A). LGIN, HGIN, and invasive ESCC can be visualized as Lugol-voiding lesions (LVLs) by Lugol chromoendoscopy.45 The staining pattern of LVLs varies from an absence of LVLs to numerous irregularly shaped multinucleated LVLs (Figure 3B). ESCC occurs synchronously and/or metachronously in conjunction with head and neck SCC. This association can be explained by the phenomenon of field cancerization.46 A polyclonal mutation in p53 is believed to be a mechanism
Figure 1. The source of acetaldehyde. The esophageal epithelium is exposed to carcinogenic acetaldehyde via (1) intrinsic and (2) extrinsic pathways. (1) Intrinsic acetaldehyde derived from ethanol metabolism in the liver. Ethanol is absorbed in the upper gastrointestinal tract and is metabolized to acetaldehyde by ADH1B in the liver. Subsequently, acetaldehyde is degraded to acetic acid by ALDH2. Acetaldehyde that exceeds its degrading activity in the liver circulates throughout the entire body, including the esophagus, lungs, and salivary glands. (2) Extrinsic acetaldehyde derived from alcoholic beverages and foods (a) and that produced by oral microflora (b). Extrinsic acetaldehyde is believed to be associated with direct exposure to the esophageal epithelium.
involved in field cancerization. A polymorphism in ALDH2 has also been related to the occurrence of multiple LVLs and multiple SCCs in the esophagus and the head and neck. Again, acetaldehyde is considered to have a key role in field cancerization of the squamous epithelium.

Pathology

In histopathologic analyses, ESCC is defined based on mitotic activity, nuclear atypia, and degree of squamous differentiation. Consequently, ESCC is classified as well-differentiated SCC, moderately differentiated SCC, poorly differentiated SCC, and undifferentiated SCC (World Health Organization classifications). The histopathologic grade is incorporated into the TNM classification as a factor (G categories: G1, well differentiated; G2, moderately differentiated; G3, poorly differentiated; G4, undifferentiated) that is used to determine prognosis. Involucrin is expressed inwell-differentiated ESCC tumor nests but not in poorly differentiated ESCC, and it is considered to be a marker of the differentiation grade of ESCC tissues. Because inhibition of squamous differentiation promotes tumor development in xenograft models, squamous differentiation might affect the malignant potential of individual ESCC cells.

Genetics

Genes that Regulate the Cell Cycle or Differentiation

Analyses of comprehensive mutational catalogues using high-throughput sequencing technologies have revealed widespread genomic alterations in ESCC (Table 1). The first large-scale comprehensive analysis, which was conducted using whole genome sequencing, whole exome sequencing, and array-based comparative genomic hybridization, showed that more than 83% of ESCCs contained a somatic mutation in TP53. The highest frequency of mutation was in TP53 (59%–93% of all patients), confirmed by other large-scale whole genome or whole exome sequencing analyses. Mutation of TP53 has therefore been proposed as a key factor in the development of ESCC.

Mutations in other cell genes that regulate the cell cycle (CDKN2A [encodes p16], RB1, NFE2L2, CHEK1, and CHEK2) or differentiation (NOTCH1 and NOTCH3) have been detected in 2% to 10% of ESCCs. Moreover, many genes that regulate the cell cycle are also amplified in ESCCs (CCND1 in 46.4% of cases, CDK4/CDK6 in 23.6% of cases, and MDM2 in 5.7% of cases), implicating them in the development of ESCC.

Epidermal Growth Factor Receptor Signaling Pathway

Epidermal growth factor receptor (EGFR) is overexpressed in 59.6% to 76% of ESCCs and is associated with a poor prognosis. Large-scale sequencing studies have shown EGFR to be overexpressed (in 68% of tumors) and/or amplified (in 11%–24%) but not frequently mutated (in 0–1.8% of ESCCs). Moreover, 78.6% of ESCCs have mutations and/or amplifications in factors downstream of EGFR, such as RAS and AKT pathways. EGFR signaling pathways are considered to be involved in the development of ESCC.

Somatic Mutational Signature

G:C→A:T transitions and G:C→C:G transversions are frequently detected in ESCC cells. Interestingly, the G:C→A:T transition is a typical mutation signature induced by acetaldehyde. Another mutation frequently detected in ESCC is that of cytosine in TpCpX trinucleotides. This pattern corresponds to the mutation signature that can be induced by members of the APOBEC family, which might reflect the increased APOBEC activity observed in ESCC cells.

Epigenetic Factors

Epigenetic alterations such as DNA methylation, histone modification, and loss of genome imprinting are involved in the development of ESCC, as for other neoplasms. For instance, hypermethylation of promoter regions of APC, RB1, and CDKN2A has been detected in ESCCs. Methylation of CDKN2A, whose product p16 regulates RB1, is associated
with p53 overexpression. These cycle-regulatory pathways appear to interact to promote the development of ESCC.

Mutations in genes that regulate histone modification have been observed in approximately 63% of ESCCs (mutations in MLL2 detected in 19%, in EP300 in 10%, in MLL3 in 6%, and in CREBBP in 6%). In addition, members of the SWI/SNF complex, which is involved in epigenetic regulation, are mutated in ESCCs (mutations in ARID detected in 5%, in PBRM1 detected in 5%, and in ARID1A detected in 1%).

**Hereditary ESCC**

Tylosis esophageal cancer, also known as Howel–Evans syndrome, is a rare familial cancer syndrome inherited in an autosomal dominant manner and characterized by hyperkeratosis of the palms or soles. People with this syndrome have a high risk of esophageal cancer. More than 90% of patients with this syndrome develop ESCC by the time they are 65 years old. Missense mutation of rhomboid family member gene 2 (RHBDF2), which is involved in the activation of EGFR signaling, has been found to cause this syndrome.

**Polymorphisms**

Polymorphisms in genes such as TP53, MDM2, CASP8, and COX2 are associated with the risk of ESCC. Recent genome-wide association study–based analyses of patients with ESCC detected several single nucleotide polymorphisms that appear to be risk factors for ESCC. These include rs4135113, which regulates nonsynonymous coding in base excision repair; rs1800450, which regulates monosaccharide binding; and rs3769823, which regulates CASP8. These single nucleotide polymorphisms are considered to be associated with susceptibility to ESCC. Moreover, an integration analysis of 3 genome-wide association studies, comprising more than 9000 cases of ESCC, associated the single nucleotide polymorphisms rs7447927 (located at 5q31.2) and rs1642764 (located at 17p13.1) with ESCC.

**Fanconi Anemia Pathway**

Fanconi anemia (FA) is an autosomal recessive disorder characterized by genomic instability, bone marrow failure, developmental defects, and early development of cancers such as hematologic malignancies and SCCs of the uterine cervix, head and neck, and esophagus. FA is caused by mutations in genes that regulate the FA pathway, which
controls replicon-dependent removal of interstrand DNA cross-links. The FA pathway is studied during research on DNA repair, cancer progression, and protein ubiquitination in response to genotoxic insults.\(^8\) FANCD2, a factor in the FA pathway, counteracts acetaldehyde-induced genotoxicity in mice.\(^8\) Moreover, germline mutations in FANCD1 (BRCA2) are also found in patients with a familial history of ESCC.\(^8\) These results indicate that the role of the FA pathway in ESCC caused by acetaldehyde-mediated DNA damage.

**HPV**

HPV promotes carcinogenesis through the action of oncoproteins E6 and E7, which target numerous cellular pathways, including the inactivation of p53 and retinoblastoma protein.\(^8\) HPV infection is associated with tumorigenesis in cervical cancer as well as in head and neck SCC.\(^8\) However, a relationship between HPV infection and development of ESCC has not been observed consistently.\(^8\) A comprehensive genetic analysis found no correlation between HPV infection and ESCC.\(^8\) In contrast, a meta-analysis reported the prevalence of HPV in patients with ESCC to be as high as 32.2%.\(^8\) Africa and Asia have the highest prevalence of HPV; specific Chinese provinces with a high prevalence of HPV have a particularly high incidence of ESCC.\(^8\) There is controversy over whether HPV infection is associated with the development of ESCC.

### Mechanisms of Invasion and Metastasis

A recent study associated amplification of the eukaryotic initiation factor 5A2 gene (eIF5A2) with ESCC invasiveness and metastasis of ESCC via hypoxia.\(^8\) eIF5A2 could promote the epithelial-to-mesenchymal transition and increase the migratory and invasive capacities of esophageal cells.\(^8\)

A 3-dimensional organotypic culture system can be used to investigate the mechanisms of invasion based on interactions between the epithelium and stroma.\(^8\) Researchers have used this model to identify several genes and signaling pathways that regulate esophageal cell invasion. Over-expression of EGFR and inactivation of p53 in esophageal epithelial cells expand a cell subpopulation via upregulation of zinc finger E-box binding transcription factors that can undergo the epithelial-to-mesenchymal transition.\(^8\) These cells have an invasive phenotype in the stromal matrix, accompanied by increases in activation of the RTK cMET.\(^3\)

Moreover, microarray analysis of RNA extracted by laser capture microdissection from invading cells grown in 3-dimensional organotypic culture, compared with non-invading cells, identified several genes that might facilitate tumor invasion. These genes include insulin-like growth factor–binding protein 3,\(^3\) peristin,\(^4\) and WNT10A.\(^5\) Furthermore, stromal fibroblasts regulate the invasive activities of ESCC cells; fibroblast-secreted hepatocyte growth factor, the ligand of c-MET, helps create an environment conducive to tumor invasion.\(^5\)

### Diagnosis and Staging

Imaging modalities such as endoscopy, endoscopic ultrasonography (EUS), esophagography, computed tomography (CT), and \(^{18}\)F-fluorodeoxyglucose positron emission tomography (FDG-PET) are used in the diagnosis and staging of ESCC. Endoscopy is the most sensitive modality for the detection and diagnosis of esophageal neoplasia. The use of endoscopic screening and treatment has contributed to reductions in ESCC-associated mortality.\(^6\) EUS, esophagography, CT, and FDG-PET are used to assess the depth of invasion in the esophageal wall, length of tumors, direct invasion to adjacent organs, and lymph node and distant metastasis.\(^6\)

#### Endoscopy

Advanced ESCC is found as a protruding mass or a depressed ulcer. Superficial ESCC is often difficult to identify because of minimal macroscopic and color changes, but it is usually observed as an uneven surface with a thin white coating or a reddish color change on the mucosal surface.\(^8\)

Iodine has an affinity for glycogen in the nonkeratinized squamous epithelium and stains the normal epithelium a mahogany brown color; the cancerous epithelium is devoid of staining because it lacks glycogen.\(^9\) Lugol solution, which contains iodine, was first introduced as Schiller test for the detection of SCC in the uterine cervix\(^9\) and is now used in

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**Table 1. Recent Results of Comprehensive Analyses of Genetic Alterations Using High-Throughput Sequencing**

<table>
<thead>
<tr>
<th>No. of patients (method)</th>
<th>Ethnic group</th>
<th>Genomic alterations</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>17 (WGS), 71 (WES), 123 (CGH)</td>
<td>Chinese</td>
<td>TP53, RB1, CDKN2A (p16), NFE2L2, CHEK1, CHEK2, PIK3CA, NOTCH1, NOTCH3, CCND1, CDK4/CDK6, MDM2, EGFR</td>
<td>Song et al(^6)</td>
</tr>
<tr>
<td>20 (WES), 119 (targeted seq), 4 (RNA seq)</td>
<td>Chinese</td>
<td>TP53, RB1, CDKN2A (p16), NFE2L2, PIK3CA, NOTCH1, NOTCH3, ARID2, EGFR, RTK/RAS signaling pathways</td>
<td>Lin et al(^6)</td>
</tr>
<tr>
<td>118 (WES)</td>
<td>Chinese</td>
<td>TP53, RB1, CDKN2A (p16), NFE2L2, CCND1, EGFR, histone-modifying genes (MLL2, MLL3, CREBBP)</td>
<td>Gao et al(^6)</td>
</tr>
<tr>
<td>14 (WGS), 90 (WES)</td>
<td>Chinese</td>
<td>TP53, CDKN2A (p16), PIK3CA, and mutational signature</td>
<td>Zhang et al(^6)</td>
</tr>
</tbody>
</table>

WGS, whole-genome sequencing; WES, whole-exome sequencing; CGH, comparative genomic hybridization; seq, sequencing.
Magnifying NBI clearly visualizes microvascular structures (intrapapillary capillary loops [IPCLs]) of the squamous epithelium and cancerous lesions (Figure 5E and F) and is useful for identifying cancerous lesions by the changes in IPCLs. A prospective, randomized, controlled study reported a higher detection rate of superficial SCC in the head and neck and the esophagus for NBI compared with conventional white light imaging (97% vs 55%). NBI detects superficial cancer in the head and neck with 100% sensitivity, 78.6% specificity, 83.3% positive predictive value, 100% negative predictive value, and 86.7% accuracy. NBI detects cancer of the esophagus with 97.2% sensitivity, 42.1% specificity, 90.4% positive predictive value, 72.8% negative predictive value, and 88.9% accuracy.

Staging

Staging of the tumor is important to determine the therapeutic strategy. The Union for International Cancer Control TNM classification (7th edition, 2009) is used widely to assess the anatomic extent of ESCC. The T stage is defined by the extent of the primary tumor and is classified as one of 5 categories: Tis (carcinoma in situ or high-grade dysplasia; eg, intraepithelial tumor), T1 (T1a, tumor invading the lamina propria or muscularis mucosa; T1b, tumor invading the submucosa), T2 (tumor invading the muscularis propria), T3 (tumor invading the adventitia), and T4 (tumor invading the adjacent structures: T4a, tumor invading the pleura, pericardium, or diaphragm; T4b, tumor invading other adjacent structures such as the aorta, vertebral body, or trachea). Endoscopy, EUS, esophagography, and CT are used for T staging of tumors.

Macroscopic findings on endoscopy are useful for the assessment of tumor depth of superficial ESCC. According to the Japanese Classification of Esophageal Cancer, superficial (type 0) cancers are classified into types 0-I (superficial and protruding type), 0-II (superficial and flat type: a lesion without definite protrusion or depression), and 0-III (superficial and excavated type). 0-I lesions and 0-III lesions are believed to have high probabilities (94.7% and 100%, respectively) of invading up to the submucosal layers (T1b stage). Most 0-IIa (slightly elevated type) and 0-IIb (flat type) tumors are Tis or T1a cancers, whereas 0-Iic (slightly depressed type) tumors are widely distributed from Tis to T1b cancers.

The depth of cancer invasion is described according to the IPCL classification pattern by NBI with magnification. Magnifying NBI can differentiate intramucosal cancers (Tis or T1a) from submucosal (T1b) cancers with high levels of sensitivity (78%) and specificity (95%). Visualization of the microvascular structure of the tumors by magnifying NBI is therefore a useful tool for the differential diagnosis of T1a from T1b cancers.

Esophagography is used for evaluating the shape, length, and location of tumor. EUS and CT are also used for evaluating the T stage of ESCC. EUS can identify the distinct tissue layers of the esophageal wall, and a tumor can be detected as a low echoic mass. EUS using a 20- to 30-MHz mini-probe is useful for diagnosing the tumor depth in superficial ESCC.
As a limitation, EUS is unreliable for the staging of tumors with stenotic lesions or after neoadjuvant chemotherapy. CT also has limitations for determining the exact depth of a tumor within the esophageal wall and thus cannot differentiate between T1, T2, and T3, although CT is useful for discriminating T3 and T4 regions. The accuracy of CT for T4b staging in terms of aortic and tracheobronchial invasion is approximately 80%. When tracheobronchial invasion is suspected, bronchoscopy and biopsy should be considered to confirm the diagnosis.

N-stage tumors (spread to regional lymph nodes) are defined by the number of involved regional nodes, including celiac axis nodes and cervical paraesophageal nodes but not supraclavicular nodes. The categories are N0 (no regional lymph node metastasis), N1 (metastasis in 1–2 regional lymph nodes), N2 (metastasis in 3–6 regional lymph nodes), and N3 (metastasis in ≥7 regional lymph nodes). EUS and CT are used to determine the presence of involved regional lymph nodes in patients with ESCC. In a meta-analysis to compare the diagnostic performance of EUS and CT in N staging, the sensitivity and specificity values for EUS were 80% and 70%, respectively; those of CT were 50% and 83%, respectively.

M-stage tumors are classified as M0 (no distant metastasis) and M1 (distant metastasis). Contrast-enhanced CT is the most commonly used imaging modality to detect distant metastasis, and masses in the liver larger than 1 cm are likely to be hepatic metastatic lesions. The sensitivity of contrast-enhanced CT for detecting metastasis is as high as 90%. FDG-PET is of value in the detection of distant metastasis at the initial staging of esophageal cancer as well as in the assessment of the response to induction chemotherapy. However, FDG-PET has a low sensitivity for initial nodal staging, and it is unclear whether FDG-PET is useful in the assessment of recurrence after surgery. Recently, coregistration of FDG-PET and CT using a combined system has been shown to be of additional value for image interpretation using both modalities. This dual-imaging modality allows clinicians to evaluate tumors functionally and structurally, improves the accuracy of FDG-PET imaging in esophageal cancer, and provides useful data of diagnostic and therapeutic significance for clinical management.

**Treatment**

There are many different approaches to the treatment of patients with ESCC, including endoscopic therapy, surgery, chemotherapy, and radiotherapy. To facilitate the standard practice of therapeutics based on the principle of evidence-based medicine, updated guidelines for esophageal cancers are published by the National Comprehensive Cancer Network. The Japan Esophageal Society has also edited guidelines for the diagnosis and treatment of esophageal carcinoma.

**Endoscopic Resection and Ablation**

Early-stage ESCC (Tis and T1a), with negligible risk of metastasis to the lymph node, can be cured by endoscopic local treatment, such as ER and/or an ablative method (eg, radiofrequency ablation or photodynamic therapy). Before endoscopic treatment, the depth of invasion, horizontal spread, and presence or absence of multifocal lesions should be characterized fully. The depth of invasion is closely associated with lymph node metastasis, and the frequency of lymph node metastasis in mucosal ESCC is reported to be 3%. However, the accuracy of pretreatment diagnosis regarding the depth of invasion using EUS and/or magnifying NBI is limited. To accurately determine the depth of invasion, diagnostic ER can be used. Endoscopic submucosal dissection gives an accurate pathological diagnosis because of the high rate of en bloc resection, irrespective of the tumor size, despite the high incidence of bleeding or perforation. ER is a standard, minimally invasive treatment for Tis and T1a ESCC. The indication of ER for T1b ESCC is controversial; a study is under way to provide the rationale to address this issue. This study concept is as follows: ER therapy alone can be effective for patients with a Tis or T1a tumor with a negative surgical margin and without lymphatic or venous invasion, but additional treatment, including surgical resection or chemoradiotherapy (CRT), should be considered for patients with a T1b tumor and/or positive surgical margin and/or positive lymphatic/venous invasion.

Endoscopic surveillance after endoscopic local therapy for early-stage ESCC should be continued because synchronous ESCC and head and neck SCC can develop, especially in patients with multiple LVls. Additional ER and/or ablation may be needed if metachronous early-stage ESCC is detected. Patients should be evaluated every 3 months in the first year after treatment and every 3 to 6 months in the second year.

**Surgery**

Surgery is used widely to obtain locoregional control and has an important role in the treatment of esophageal cancer. Transthoracic esophagectomy is one of the most invasive surgeries. Outcomes appear to be related to the incidence and management of perioperative complications and seem to be good in high-volume centers with experienced surgeons. ESCC is often accompanied by extensive metastasis to lymph nodes in the cervical, thoracic, and abdominal regions. There is controversy regarding the extent of lymph node dissection required. In particular, the advantage of a 3-field lymphadenectomy over a 2-field lymphadenectomy for ESCC is unclear. A recent meta-analysis of 13 studies revealed a marked improvement in surgical outcomes after 3-field lymphadenectomy compared with 2-field lymphadenectomy, including a higher 5-year rate of survival (hazard ratio, 0.64). On the other hand, perioperative morbidities have been noted, such as a higher prevalence of anastomotic leakage and a similar prevalence of pulmonary complications and postoperative vocal cord palsy. The addition of cervical lymph node dissection therefore improves the long-term outcomes of patients compared with only 2-field lymphadenectomy, especially for patients with thoracic esophageal cancer with positive lymph nodes.
However, 3-field lymphadenectomy is regarded to be the more invasive procedure.\textsuperscript{125}

To overcome this issue, minimally invasive surgery has been developed. Magnifying the view from the thoracoscope makes it possible to perform accurate lymph node dissection; this technique has been applied to patients with stage I, II, and III (excluding T4) esophageal cancers. Some evidence is available on the short-term benefits (shorter hospital stays as well as lower rates of respiratory complications and total morbidity) of minimally invasive surgery compared with open esophagectomy.\textsuperscript{126,127} Moreover, the rate of 5-year survival after minimally invasive esophagectomy is similar to that for open esophagectomy.\textsuperscript{128} A prospective randomized trial is under way in Japan to prove the non-inferiority of minimally invasive thoracoscopic esophagectomy to open esophagectomy for thoracic esophageal cancers.

**Neoadjuvant and Adjuvant Therapy**

Neoadjuvant chemotherapy (in the United Kingdom) or neoadjuvant CRT (in the United States and France) is performed as standard treatment for locally advanced ESCC.\textsuperscript{129,130} However, clinical trials conducted in Europe and the United States have included a low percentage of patients with SCC (<50%). Randomized multicenter trials of patients with stage I/II ESCC that compared neoadjuvant CRT (cisplatin + 37-Gy radiation) followed by surgery with surgery alone showed that neoadjuvant CRT did not increase overall survival.\textsuperscript{131} However, a recently updated meta-analysis (24 trials, 4188 patients, 3500 events, 65% of patients with SCC) provided strong evidence that neoadjuvant CRT and chemotherapy increase survival versus surgery alone (hazard ratios of 0.78 and 0.87, respectively) in patients with esophageal cancer, although a clear advantage of neoadjuvant CRT over neoadjuvant chemotherapy has not been shown.\textsuperscript{132} Of note, when limited to the patients with ESCC included in this meta-analysis, neoadjuvant CRT significantly increased the chances of survival versus surgery alone (hazard ratio, 0.80) but not versus neoadjuvant chemotherapy (hazard ratio, 0.92).\textsuperscript{132}

In Japan, neoadjuvant chemotherapy with cisplatin plus 5-fluorouracil (5-FU) is the standard treatment regimen for locally advanced (stage II or III, except T4) ESCC.\textsuperscript{133} This neoadjuvant chemotherapy increases overall survival and is regarded as the standard treatment for patients with stage II or III ESCC (55% survive for 5 years).\textsuperscript{134} Based on this background, another intensive neoadjuvant chemotherapeutic regimen with docetaxel and cisplatin plus 5-FU (DCF) has been proposed.\textsuperscript{135} A phase 3 trial is under way to investigate whether DCF is superior to cisplatin plus 5-FU and whether cisplatin plus 5-FU with CRT is superior to cisplatin plus 5-FU alone.\textsuperscript{136}

Interestingly, a German trial reported that the addition of surgery did not prolong survival of patients with locally advanced thoracic ESCC (T3-N0-1M0) who responded to neoadjuvant chemotherapy followed by CRT.\textsuperscript{137} In this trial, a tumor response to neoadjuvant chemotherapy was the single prognostic factor for overall survival.\textsuperscript{139} A trial in France also reported that the addition of surgery did not prolong survival of patients with locally advanced thoracic esophageal cancer (T3-N0-1M0; 89% of patients with SCC) who responded to induction CRT.\textsuperscript{138} These results indicate that good responses to neoadjuvant chemotherapy or CRT may not require additional surgery.

**Definitive CRT**

Because CRT has been reported to be superior to radiotherapy alone for the treatment of patients with locally advanced esophageal cancer,\textsuperscript{139} the concept of definitive CRT has emerged. At first, definitive CRT was applied to inoperable locally advanced (T4) ESCC. Even for T4 ESCC, definitive CRT can produce complete remission in 15% to 33% of patients (median survival time of approximately 10 months).\textsuperscript{140,141}

A clinical trial of definitive CRT, which included patients with stage II or III ESCC, reported a complete response in 68% of patients and that 37% survived for 5 years.\textsuperscript{142} This result was inferior to that of neoadjuvant chemotherapy followed by surgery (5-year rate of survival, 55%).\textsuperscript{134} Definitive CRT for stage II or III ESCC is therefore a nonsurgical treatment option.

CRT has some problems. First, CRT fails in more than 40% of patients\textsuperscript{143}; the prognosis is poor for these patients, with more than 50% mortality within 6 months.\textsuperscript{144} In such cases, salvage treatment is needed. Second, approximately 10% of patients with complete responses after treatment with 60-Gy CRT experience late-stage, high-grade toxicity, including pericarditis, pleural effusion, and radiation pneumonitis.\textsuperscript{145} A dose-escalation study of radiation was conducted to improve local control.\textsuperscript{146} However, no significant difference was observed in local/regional control of patients (85% of patients with SCC) treated with cisplatin plus 5-FU in the 50.4-Gy group versus the 64.8-Gy group.\textsuperscript{146} Based on this evidence, the standard radiation dose of definitive CRT for ESCC is currently 50 to 50.4 Gy at 1.8 to 2 Gy per fraction. In addition, the use of the multiple-field technique to reduce the volume of the heart within the radiation field is recommended to prevent late-stage cardiac toxicity.

**Salvage Therapy**

The diagnosis of local recurrence after CRT should be conducted promptly and precisely. The median time to local recurrence after CRT is 9 months.\textsuperscript{147} Early recurrent tumors that have achieved complete remission after CRT typically show a submucosal tumor-like appearance (Figure 6).\textsuperscript{148}

Salvage therapies, including ER and surgery, are considered for residual or recurrent tumors after definitive CRT.\textsuperscript{149} Salvage ER has been tested on residual and recurrent tumors at primary sites after definitive CRT and produced positive long-term results without severe complications in patients with superficial failed lesions.\textsuperscript{150} Recently, photodynamic therapy has been shown to be a curative option for patients with local failure limited to the submucosal layer.\textsuperscript{151} Salvage surgery is aimed at curative resection for those tumors. However, only patients with T1N0 or T2N0 ESCC survive for long periods.\textsuperscript{144} Salvage
surgery has a greater risk of respiratory complications and anastomotic leakage than planned surgery.\textsuperscript{144,152} This risk is associated with fibrous changes in the mediastinum after radiation and difficulties in the anastomosis as a result of the irradiated gastric tubes.\textsuperscript{144}

**Chemotherapy for Unresectable Locally Advanced or Metastatic ESCC**

Combinations of cisplatin and 5-FU are commonly used in chemotherapy for patients with unresectable locally advanced or metastatic ESCC; this treatment is believed to be better than the best supportive care.\textsuperscript{153} Taxanes (docetaxel and paclitaxel) have been reported to be effective as single-agent chemotherapy, with response rates of 20\% to 34\%.\textsuperscript{154,155} Recently, the 3-drug regimen with DCF was reported to have a 62\% response rate.\textsuperscript{156} Based on this result, a randomized controlled phase 3 study is under way to compare DCF with cisplatin plus 5-FU in patients with metastatic or recurrent ESCC.\textsuperscript{157}

**Targeted Therapy**

EGFR is one of the most investigated molecular targets in the field of SCC. Cetuximab, a mouse-human monoclonal immunoglobulin G1 against EGFR, is effective against head and neck SCC in combination with radiotherapy.\textsuperscript{158} However, addition of cetuximab to CRT did not yield a significant survival benefit for esophageal cancer, including ESCC, in a phase 3 study.\textsuperscript{159} A phase 3 study of gefitinib, a tyrosine kinase EGFR inhibitor, in patients with ESCC did not show an increase in overall survival.\textsuperscript{160} There is therefore little evidence to support EGFR-targeting therapies for ESCC.

Recently, strategies have been developed to target the programmed cell death protein 1 signaling pathway (PD1-PDL1).\textsuperscript{161} Patients whose ESCC was positive for PD1 signaling (43.9\%) had significantly poorer outcomes than patients whose ESCC did not have activation of this signaling pathway.\textsuperscript{162} On the basis of these data, a clinical trial is under way to target the PD1-PDL1 pathway in patients with ESCC who have not responded to standard chemotherapy.

**Palliative Care**

Some patients with advanced ESCC experience dysphagia and malnutrition because of esophageal stenosis and aspiration caused by a fistula.\textsuperscript{120} To improve these conditions, palliative care is strongly recommended.

Dysphagia and malnutrition decrease quality of life in patients with advanced ESCC.\textsuperscript{163} Radiation therapy palliates dysphagia for several months, but it takes 4 to 6 weeks after treatment for symptoms to improve.\textsuperscript{164} Instead, a self-expanding metal stent (SEMS) is a safe and effective treatment for palliation of dysphagia in patients with ESCC.\textsuperscript{165} Compared with uncovered SEMSs, covered SEMSs prevent tumor growth (53\% vs 100\%) and reduce restenosis (8\% vs 37\%).\textsuperscript{166} Biodegradable stents have been developed, although their application is limited because of unknown long-term efficacy.\textsuperscript{167} In patients with prior CRT, SEMSs can cause life-threatening complications.\textsuperscript{168}
Placement of a SEMS is also effective for closing malignant esophagorespiratory fistulas. SEMS were shown to seal the esophagorespiratory fistula successfully in 80% (49/61) of patients. Patients with successful closure with SEMSs survived longer than those with unsuccessful closure (15.1 vs 6.2 weeks).

Future Directions

Alcohol, acetaldehyde associated with alcoholic beverages, and tobacco are esophageal carcinogens that contribute to the development of ESCC. Polymorphisms in ALDH2 also contribute to the development of ESCC, particularly in Asian people. In addition, external and internal risk factors together contribute to the development of ESCC. An effort is needed to develop preventive strategies as the next step. Advances in early detection should encourage the development of effective screening systems for high-risk people. Effective detection of early ESCC allows for the use of minimally invasive treatments such as ER. Field cancerization should be considered for intensive surveillance in successfully treated patients. However, for advanced ESCC, the treatment outcomes leave room for improvement. Further research is needed to establish techniques for less invasive surgery; more effective chemotherapy and radiotherapy are required to increase patient survival. Precision medicine based on genomic data could lead to new methods for prevention, diagnosis, and treatment of ESCC.

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