

Japanese eCura system

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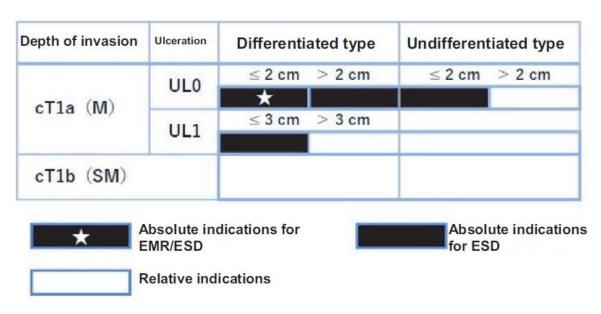


Figure 1 Classification of indications according to tumor-related factors. cT1a (M), intramucosal cancer (preoperative diagnosis), cT1b (SM), submucosally invasive cancer (preoperative diagnosis). UL, finding of ulceration (or ulcer scar); UL0, absence of ulceration or ulcer scar; UL1, presence of ulceration or ulcer scar.

Post-ESD curability criteria

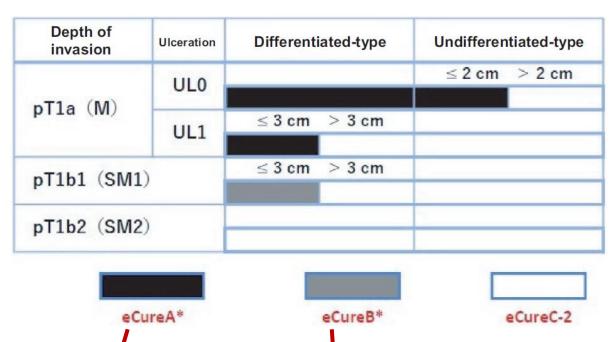


Figure 2 Evaluation of curability according to tumor-related factors. *, Confined to *en bloc* resection and HMO, VMO, LyO, and VO. pT1a (M), intramucosal cancer (histopathological diagnosis); rT1b (SM), submucosally invasive cancer (histopathological diagnosis). UL, finding of ulceration (or ulcer scar); ULO, absence of ulceration or ulcer scar; UL1, presence of ulceration or ulcer scar.

Curative resection

Expanded curative resection

eCuraC-1 vs eCuraC-2

Endoscopic curability C

This level of curability corresponds to the concept of noncurative resection described in the Guidelines for ESD and EMR for EGC (first edition). When a lesion meets neither of the above-mentioned eCuraA and B conditions, it is considered eCuraC, which has a likelihood of remnant tumor. When eCuraC lesions are differentiated-type lesions and fulfill other criteria to be classified into either eCuraA or eCuraB but was either not resected *en bloc* or had positive HM, they are considered eCuraC-1. All other eCuraC lesions are considered eCuraC-2.

Management by eCura system

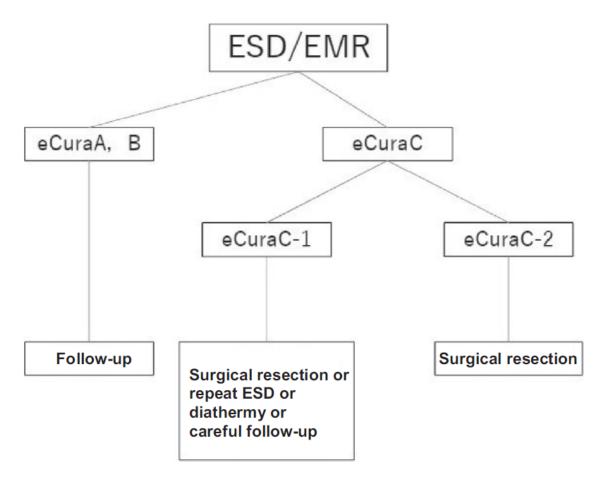


Figure 3 Therapeutic flowchart following endoscopic submucosal dissection (ESD) or endoscopic mucosal resection (EMR).

Japanese multicenter study

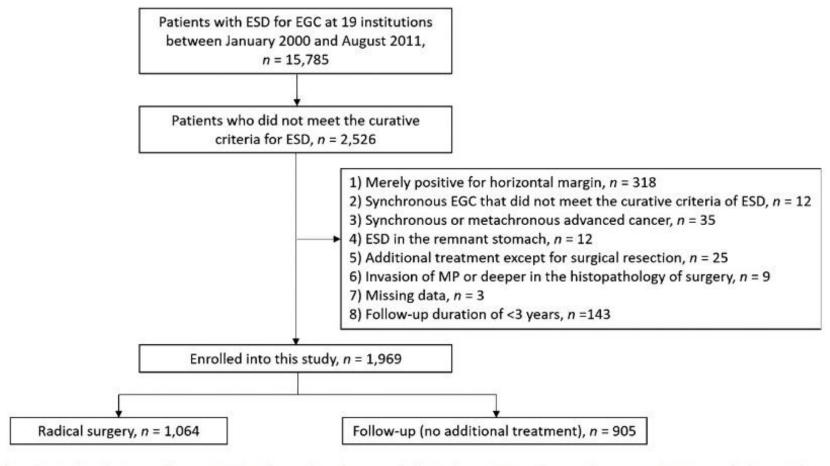
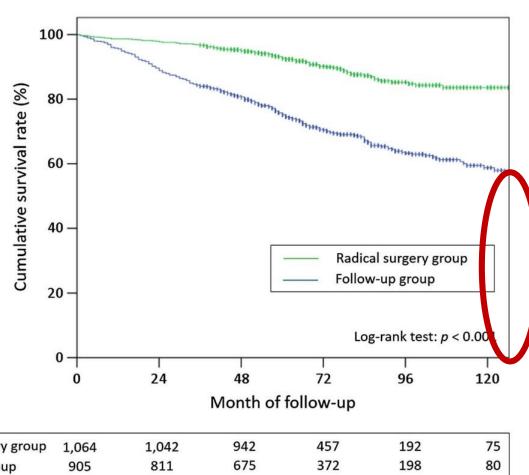


Fig. 1 Flowchart of patient enrollment. ESD endoscopic submucosal dissection, EGC early gastric cancer, MP muscularis propria

그러나 모두 사망하는 것은 아니다.

Fig. 2 OS rate for patients who did not meet the current curative criteria for ESD of EGC. The 3-year OS rate was 96.7 % in the radical surgery group and 84.0 % in the follow-up group, which was significantly different (p < 0.001). OS overall survival. ESD endoscopic submucosal dissection, EGC early gastric cancer



Number at risk

Radical surgery group	1,064	1,042	942	457	192	75
Follow-up group	905	811	675	372	198	80

어느 정도 예측할 수 있었던 바이다.

- LN metastasis in lesions without LV infiltration

Category		Rate of lymph node metastasis	
1)	>3 cm in long diameter, differentiated type, pT1a, UL1	3.0% (95% CI: 1.2-6.2%)	
2)	>3 cm, differentiated type, pT1b1 (SM1)	2.6% (95% CI: 0.3-9.0%)	
3)	>2 cm, undifferentiated type, pT1a, UL0	2.8% (95% CI: 1.0-6.0%)	
4)	≤2 cm, undifferentiated type, pT1a, UL1	2.9% (95% CI: 1.2-5.7%)	
5)	>2 cm, undifferentiated type, pT1a, UL1	5.9% (95% CI: 4.3-7.9%)	
6)	Undifferentiated type, pT1b1 (SM1)	10.6% (95% CI: 5.0-19.2%)	

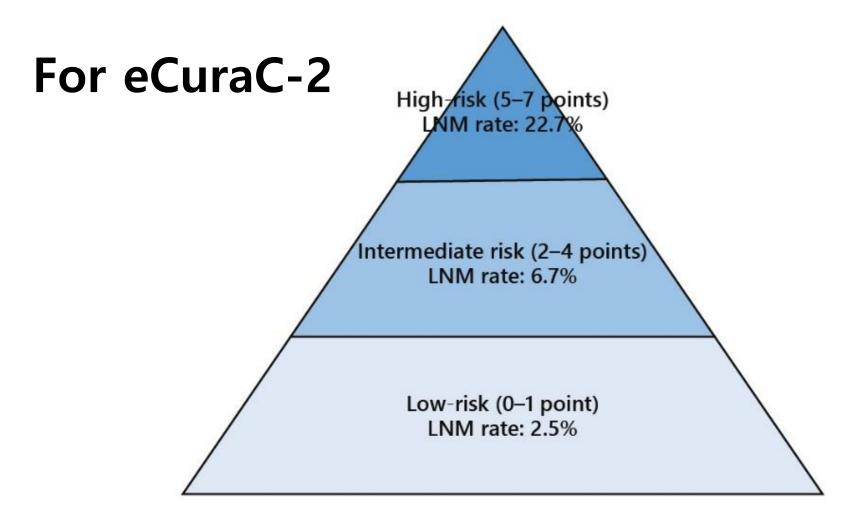
0			- 2 am	
			≥ Z CIII	> 2 cm
3754111				3)
1	≤ 3 cm	> 3 cm		
· _		1)	4)	5)
1, 3	≤ 3 cm	> 3 cm		
		2)	6)	
			1) ≤ 3 cm > 3 cm	1 1) 4) ≤ 3 cm > 3 cm



Table 2. Multivariate logistic regression analysis of risk factors for LNM in the development cohort and scoring system No. of patients No. of LNMs 95% CI Points^b OR P value β regression coefficient Tumor size 2.03 0.003 0.70 >30 mm 479 53 1.28-3.14 ≤30 mm 622 41 Reference 1 Tumor depth SM₂ 197 30 1.68 0.97-2.92 0.065 0.52 M/SM1 904 64 Reference 1 Histopathological type 1.22 0.56 0.20 Undifferentiated 701 73 0.62 - 2.41Differentiated 400 21 1 Reference Lymphatic invasion Positive 3.99 2.43-6.55 443 69 < 0.001 1.38 25 Negative 658 1 Reference Venous invasion Positive 249 1.65 1.01-2.70 0.046 0.50 35 1 Negative 852 59 Reference Ulceration (scar) Presence 285 21 0.98 0.57-1.69 0.95 -0.016816 73 Reference Absence 1 Vertical margin 198 1.81 1.10-3.00 0.020 0.60 Positive 30 Otherwise 903 64 1 Reference

LN metastasis by the score

Total score	Rate of lymph node metastasis	No. of cases
0	1.6% (95% CI: 0.3–8.6%)	1/62
1	2.6% (95% CI: 1.4-4.9%)	9/341
2	4.9% (95% CI: 2.6-9.0%)	9/185
3	7.4% (95% CI: 4.2-12.8%)	11/148
4	8.3% (95% CI: 4.7-14.3%)	11/132
5	19.9% (95% CI: 14.1-27.2%)	28/141
6	27.3% (95% CI: 18.6-38.1%)	21/77
7	26.7% (95% CI: 10.9-52.0%)	4/15



3 points: Lymphatic invasion 1 point: Tumor size >30 mm

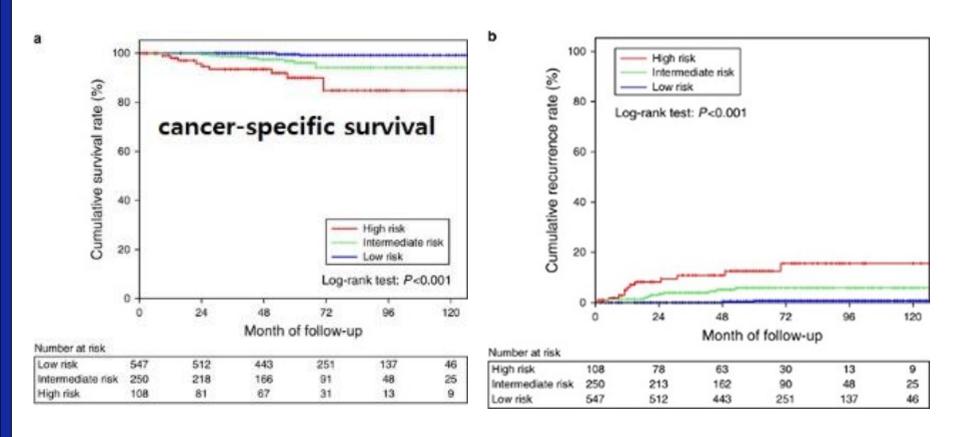
Positive vertical margin pT1b-SM2

Vascular invasion

0 point: Undifferentiated-type

Ulceration (scar)

Survival and recurrence rate



미분화 혼재암은?

미분화 혼재 점막암

Endoscopic curability A: curative resection

Endoscopic resection is shown to be equal or superior to surgical resection in terms of long-term outcomes. 10,11 When the lesion is resected en bloc, the following conditions: (i) predominantly differentiated type, pT1a, UL0, HM0 VM0, Ly0, V0, regardless of size; (ii) long diameter ≤2 cm, predominantly undifferentiated type, pT1a, UL0, HM0, VM0, Ly0, V0; or (iii) long diameter ≤3 cm, predominantly differentiated type, pT1a, UL1, HM0, VM0, Ly0, V0, are considered for endoscopic curability A (eCuraA). 62,63 However, evidence is lacking for cases of differentiated-type cancers with undifferentiated components. The above-mentioned type (1) lesions with the undifferentiated components measuring >2 cm in long diameter are defined as endoscopic curability C (eCuraC)-2 (see the measuring method in Fig. 6).

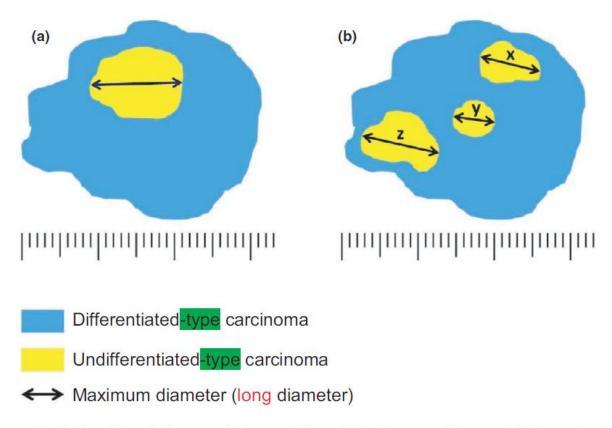


Figure 6 Measurement of the size of the coexisting undifferentiated-type carcinoma. (a) Reconstruct the area of the undifferentiated-type carcinoma, and measure the long diameter of that area. (b) If undifferentiated-type carcinoma is present in more than one area, measure the long diameters (x, y, z) of all these areas and record the sum of these values.

미분화 혼재 점막하암

Endoscopic curability B

Although no sufficient long-term results have yet been obtained, curability can be expected. When the lesion is resected en bloc, is ≤ 3 cm in long diameter, predominantly of the differentiated type, and satisfies the following criteria: pT1b1(SM1) (within <500 µm from the muscularis mucosae), HM0, VM0, Ly0, and V0, it is considered endoscopic curability B (eCuraB).⁶⁴ However, the lesion is considered eCuraC-2 (see the measuring method in Fig. 6) if undifferentiated components are present in the submucosally invasive part of the lesion.