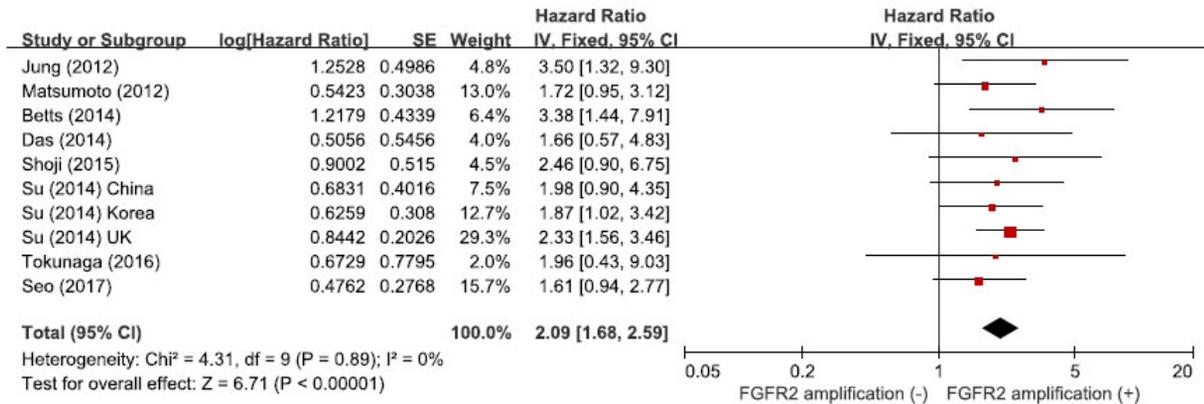


Prognostic impact of FGFR2 amplification in gastric cancer: a meta-analysis and systemic review

대한민국 한림대학교 강남성심병원 혈액종양내과

*이경무, 김형수, 김정환

Background/Aims: Fibroblast growth factor receptor-2 (FGFR2) amplification has been reported in up to 15% of patients with gastric cancer (GC). However, the prognostic impact of FGFR2 amplification has not been consistent among studies. **Methods:** We conducted this meta-analysis to evaluate the prognostic significance of FGFR2 amplification in patients with GC. A systematic computerized search of the electronic databases including PubMed, PMC, EMBASE, Google scholar, and Cochrane Library (up to June 2018) was performed. **Results:** From eight studies, 2,377 patients were included in the pooled analysis of hazard ratios (HRs) with 95% confidence intervals (CIs) for overall survival. There was no significant heterogeneity among studies ($X^2=4.32$, $P=.89$, $I^2=0\%$). Compared with patients with FGFR2-unamplified GC, patients with tumors harboring FGFR2 amplification showed significantly worse survival (HR=2.09, 95% CI: 1.68-2.59, $P<.00001$). **Conclusions:** In conclusion, this meta-analysis indicates that FGFR2 amplification is an adverse prognostic factor in patients with GC. However, the potential usefulness of FGFR2 amplification as a predictive biomarker for response to FGFR2 inhibitors remains to be investigated.



A pilot study of exercise intervention in patients with metastatic cancer

¹가천의과대학 길병원 내과, ²가천의과대학 길병원 영상의학과, ³가천의과대학 길병원 재활의학과

*이주영¹, 박주환¹, 김영생¹, 김은영², 안희경¹, 박인근¹, 박기덕³, 심선진¹, 신동복¹

Background/Aims: Skeletal muscle loss is a central component of cancer cachexia syndrome and is a poor prognostic factor in cancer patients. We investigated prevalence of sarcopenia and the feasibility of exercise intervention in patients with advanced solid cancers receiving first-line palliative chemotherapy. **Methods:** Between July 2017 and February 2018, consecutive patients with newly diagnosed with advanced solid cancer were enrolled to our prospective cohort of sarcopenia. Skeletal muscle mass was measured using both the cross-sectional area of muscle at the level of the third lumbar vertebra(L3) on baseline computed tomography and bioelectrical impedance analysis(BIA). Sarcopenia was defined as a L3 muscle index of less than 55cm²/m² for men and of less than 39cm²/m² for women. Patients were recruited to participate in a 12-week, combined resistance and aerobic exercise program consisting of supervised, hospital-based training and home-based training, during the first-line palliative chemotherapy. The primary endpoint was feasibility and safety of the exercise intervention. Skeletal muscle mass by BIA and patients' quality of life questionnaires were measured at baseline and after the intervention. **Results:** Among 76 patients enrolled, sarcopenia was present in 58 patients. 19 patients was enrolled in the exercise program, however 5 patients withdrew consent before commencement(health concern(n=2), distance to the hospital(n=1), unspecified(n=2)). The completion rate of the 12-week exercise program was 78.6%. Disease progression(n=2) was the main reason for early discontinuation. The adherence rate of the supervised exercise session was 78.1% (207/265) and there was no adverse event associated with the exercise. Among participants in the program, there was no significant change in skeletal muscle index (mean, 9.4±1.3 kg/m² vs. 9.4±1.2 kg/m², $p=0.982$). FACIT-fatigue scale non-significantly improved after the exercise intervention (mean, 35.2±10.4 vs. 38.2±9.8, $p=0.635$). **Conclusions:** Exercise interventions appear to be feasible and safe in patients with advanced solid cancer and might have a role of preventing skeletal muscle loss without fatigue exacerbation during palliative chemotherapy.

Table 1. Baseline characteristic of the study population^a

Variable ^b	Cohort ^c (n = 76) No. (%) ^d	Exercise group ^e (n=19) No. (%) ^d
Age, years ^e		
median (range) ^e	64 (30-84) ^e	60 (30-74) ^e
Male ^e	52 (68.4) ^e	10 (52.6) ^e
BMI, kg/m ² ^e		
Underweight (< 18.5) ^e	4 (5.3) ^e	0 (0) ^e
Normal (≥ 18.5, < 23.0) ^e	37 (48.7) ^e	8 (42.1) ^e
Overweight (≥ 23.0, < 25.0) ^e	21 (27.6) ^e	7 (36.8) ^e
Obese (≥ 25.0) ^e	14 (18.4) ^e	4 (21.0) ^e
Stage (TNM) ^e		
III ^e	2 (2.6) ^e	0 (0) ^e
IV ^e	74 (97.4) ^e	19 (100) ^e
ECOG PS ^e		
0 ^e	15 (19.7) ^e	7 (36.8) ^e
1 ^e	58 (76.3) ^e	12 (63.2) ^e
2 ^e	3 (4.0) ^e	0 (0) ^e
Cancer type ^e		
Lung cancer ^e	32 (42.1) ^e	9 (47.3) ^e
Gastrointestinal cancer ^e	28 (36.8) ^e	1 (5.2) ^e
Gynecologic cancer ^e	7 (9.2) ^e	5 (26.3) ^e
Genitourinary cancer ^e	1 (1.3) ^e	0 (0) ^e
Others ^e	8 (10.5) ^e	4 (21.1) ^e
Sarcopenia ^e	58 (76.3) ^e	15 (78.9) ^e

BMI, body mass index; ECOG PS, Eastern Cooperative Group Performance Status; ^e -

Table 2. Change of skeletal muscle mass and quality of life in exercise participants^a

Measure ^b	Before exercise ^c	After exercise ^c	p value ^d
Skeletal muscle mass index (kg/m ²) ^e	9.42 (±1.26) ^e	9.42 (±1.18) ^e	0.982 ^e
FACIT fatigue scale ^e	35.2 (±10.36) ^e	38.3 (±9.79) ^e	0.635 ^e

FACIT, Functional Assessment of Chronic Illness Therapy

^aResults presented as mean±SD.