

Overexpression of Romo1 is an unfavorable prognostic biomarker and a predictor of lymphatic metastasis in non-small cell lung cancer patients

Hong Jun Kim¹
 Min Jee Jo²
 Bo Ram Kim²
 Jung Lim Kim²
 Yoon A Jeong²
 Yoo Jin Na²
 Seong Hye Park²
 Suk-young Lee¹
 Dae-Hee Lee^{1,2}
 Baek-hui Kim³
 Young Do Yoo⁴
 Sang Cheul Oh^{1,2}

¹Division of Oncology and Hematology, Department of Internal Medicine, College of Medicine, Korea University, Seoul, Republic of Korea;

²Laboratory of Oncology, Graduate School of Medicine, Korea University, Seoul, Republic of Korea; ³Department of Pathology, College of Medicine, Korea University, Seoul, Republic of Korea; ⁴Laboratory of Molecular Cell Biology, Graduate School of Medicine, Korea University, Seoul, Republic of Korea

Correspondence: Sang Cheul Oh
 College of Medicine, Korea University,
 148, Gurodong-ro, Guro-gu,
 Seoul 08308, Republic of Korea
 Tel +82 2 2626 3060
 Email sachoh@korea.ac.kr

Introduction: Reactive oxygen species modulator-1 (Romo1) is a protein that modulates levels of reactive oxygen species (ROS) and has been reported to affect cancer cell invasion and proliferation via persistent inflammation. Several studies have demonstrated the clinical application of Romo1 as a prognostic marker in non-small cell lung cancer (NSCLC); however, there have been no studies investigating the mechanism by which Romo1 adversely affects the prognosis of these patients.

Methods: We examined Romo1, ROS, and vascular endothelial growth factor (VEGF) in tumor tissues immunohistochemically. We conducted survival analyses of patients who had curative resection (n=30) in accordance with clinical parameters including levels of Romo1 expression.

Results: Romo1 levels were associated with serologic inflammatory markers and high lymphatic metastatic tendencies. Significantly longer disease-free survival (68.7 vs 24.2 months, $P=0.031$) and overall survival (92.7 vs 51.6 months) were observed in the group with low Romo1 compared with high Romo1. Survival outcomes were also significantly associated with serologic inflammatory markers. Spearman's correlation analyses demonstrated significant positive correlations of Romo1 expression with VEGF-C ($P=0.008$, $R=0.478$) and ROS ($P=0.016$, $R=0.436$) in tumor samples.

Conclusion: The current study demonstrates that Romo1 induces lymphatic metastasis of NSCLC by modulating persistent inflammation and oxidative stress (ROS)/VEGF signaling. Lymphatic metastasis associated with elevated Romo1 was shown to be a key reason for unfavorable survival rates.

Keywords: non-small cell lung carcinoma, inflammation, reactive oxygen species modulator-1, vascular endothelial growth factor A, reactive oxygen species

Introduction

Lung cancer is the leading cause of cancer deaths, with an estimated 1.6 million new cases and 1.4 million deaths annually worldwide.¹ In the Republic of Korea, there are 23,000 new cases and ~16,000 deaths annually, which represents 22.6% of all cancer-related mortalities.² The World Health Organization classifies lung cancers into four groups: small cell lung cancer, non-small cell lung cancer (NSCLC) including adenocarcinoma, squamous cell carcinoma, and large cell carcinoma. Approximately half of all patients with NSCLC which account for about 80% of lung cancer diagnoses³ are initially diagnosed as having metastatic disease which is not suitable for local treatment; thus, the role of systemic chemotherapy is crucial. Despite increasing use of targeted therapies for systemic treatment including epidermal growth factor receptor,

tyrosine kinase inhibitor, and anaplastic lymphoma kinase inhibitor, 5-year survival rates of patients with NSCLC remain disappointingly low at <20%.⁴ Recently, the introduction of immune checkpoint inhibitors is promising; however, at present, the only available biomarker for their use is programmed death-ligand 1 (PD-L1).⁵ Continued research on appropriate biomarkers, and not just for immunotherapy, is required to develop new anticancer agents, expand indications for targeted agents or immune checkpoint inhibitors, and establish combination therapies required to maximize efficacy.

There is not a great deal of controversy about whether persistent inflammation is a trademark of malignancy.⁶ Chronic inflammation is also thought to be one of the barriers against CD8⁺ T-cells which are major anticancer effector cells in the immuno-oncology field.⁷ Recently, a novel protein called reactive oxygen species modulator-1 (Romo1), that induces mitochondrial production of reactive oxygen species (ROS), has been reported to affect cancer cell invasion and proliferation via persistent inflammation.^{8,9} It was first discovered in 2006 as a newly overexpressed gene in malignant tissues that had become resistant in the course of systemic chemotherapy.¹⁰ A few studies have demonstrated the clinical application of Romo1 as a prognostic marker in NSCLC.^{11,12} In another study, upregulated Romo1 was shown to be associated with poorer survival rate of patients with hepatocellular carcinoma (HCC) and the authors suggested that cancer invasiveness induced by Romo1 was the reason for poor survival.¹³ Romo1 was also shown to have potential as an adverse prognostic marker in patients with colorectal cancer (CRC), and was reported to be associated with high lymphatic metastatic tendency.¹⁴

The aim of the current study was to demonstrate the value of Romo1 as a prognostic marker, to check the correlation between Romo1 and serologic biomarkers of persistent inflammation, and to investigate the mechanism by which Romo1 affects the prognosis of patients with NSCLC who had undergone curative resection.

Methods

Study patients and specimens

We collected formalin fixed, paraffin-embedded samples from patients with NSCLC who underwent curative resection in Korea University Guro Hospital from 2007 to 2010. All patients whose tissues were used in this research provided written informed consent for their use in future research. Patients who had died within 1 month of surgical resection, or received any treatment including chemotherapy or concurrent chemoradiotherapy before surgery, were

excluded to reduce bias in the survival analysis. We also excluded three cases whose samples were not available for immunohistochemical (IHC) staining; thus, a total of 30 patients were enrolled. We collected the samples after surgery and verified pathologic staging according to the seventh edition of the American Joint Committee on Cancer (AJCC) Cancer Staging Manual. We also collected relevant clinical data by reviewing medical records retrospectively. This study was performed with permission of the Institutional Review Board of Guro Hospital and written informed consent was obtained from all patients (KUGGR-2010-057).

Immunohistochemistry for Romo1, vascular endothelial growth factor (VEGF), and ROS

Tissue microarray (TMA) recipient blocks were made by using the collected paraffin blocks as donor blocks. In each donor block, morphologically ideal areas were designated and marked on their respective hematoxylin and eosin stained slides. Tissue cores of 0.6 mm diameter from all donor blocks were taken by a cylindrical tissue puncher (MicroDigital Co., Ltd. Seoul, Korea) and transferred into the hole on the recipient paraffin blocks.

Immunohistochemistry was used to demonstrate ROMO1, VEGF-C, and VEGF-D expression in primary tumor tissues. We deparaffinized the TMA slides with xylene and incubated them with peroxidase blocking reagent (3% H₂O₂ diluted with dextrose water) for 15 minutes to eliminate endogenous peroxidase activity. Subsequently, we retrieved antigens by steaming samples at 95°C for 20 minutes and freezing them for 10 minutes. Then, we blocked the TMA slides with blocking buffer, and incubated them with antibodies for Romo1 (OriGene, Rockville, MD, USA) at a 1:150 dilution, VEGF-D (Abcam, Cambridge, UK) at a 1:50 dilution, and VEGF-C (Abnova, Taipei City, Taiwan) at a 1:25 dilution. We then treated samples with 3,3'-diaminobenzidine chromogen solution for 7 minutes, followed by counterstaining with hematoxylin and dehydration. Negative control slides without the primary antibody were included.

For determination of ROS levels, the above-constructed TMA blocks were examined using a ROS detection kit (1:100, OxyIHC Oxidative Stress Detection Kit; Merck Millipore, Billerica, MA, USA) according to the manufacturer's instructions.

All IHC staining was evaluated independently under a light microscope by two investigators (Baek-hui Kim and Hong Jun Kim) who were unaware of the clinical information. When cytoplasmic staining was observed, the stained

cells were recorded as positive. The staining intensity was graded from 0 to 3, and the staining percentage of positive cells was classified into six grades (percentage score 0–5). Results from the two evaluators were ~80% identical. When the results were discordant between the two investigators, a third party conducted an inspection. IHC score was calculated by multiplying the percentage score by the intensity score (possible range 0–15).

Serologic inflammatory biomarkers

Evaluation of serologic inflammatory biomarkers, including white blood cell (WBC), neutrophil, lymphocyte, C-reactive protein (CRP), albumin, and platelet (PLT), was performed just before patients' operations. The neutrophil–lymphocyte ratio (NLR) was calculated as the absolute neutrophil count divided by the absolute lymphocyte count; the platelet–lymphocyte ratio (PLR) was calculated as the PLT count divided by the absolute lymphocyte count. The CRP–albumin ratio was calculated as the serum CRP level (mg/L) divided by serum albumin level (g/dL).

Statistical analysis

We estimated the disease-free survival (DFS) and overall survival (OS) as clinical outcome variables. DFS was defined as the period from surgical resection until the time of the first detection of recurrence, or death from any cause. OS was defined as the period from surgical resection to death from any cause; we censored data for cases without cancer recurrence or death, at the last follow-up. Survival curves were constructed by the Kaplan–Meier method and analyzed by the log-rank test.

We categorized patients into two groups according to each clinical variable. For WBC, CRP, and PLT, patients were grouped by their upper normal values. For albumin level, patients were divided by the lower normal value. For those clinical data with no known normal range (Romo1, NLR, PLR, and CRP–albumin ratio), we performed receiver operating curve (ROC) analysis using 5-year survival rates to determine the optimal cutoff levels. We assessed the relationship between clinical parameters including serologic biomarkers with the Romo1 levels, by chi-square test. Using the Mann–Whitney *U* test, we measured the associations between Romo1 levels in two patient groups by clinical parameters. We considered a *P*-value <0.05 as statistically significant and used a Bonferroni corrected *P*-value (*P_c*) for multiple comparisons. All statistical analyses were performed using SPSS version 20.0 for Windows (IBM Corporation, Armonk, NY, USA).

Results

Patient characteristics

Patient characteristics are shown in Table 1. Among 30 patients, eight (23.3%) were female and 22 (76.7%) were male. Their median age was 65.2 years (range 44–84 years). Eighteen patients (60%) were never smokers and 12 patients (40%) were current or former smokers. Nineteen patients (63.3%) had adenocarcinoma, nine had squamous cell carcinoma (30.0%), and two had pleomorphic carcinoma (6.7%). Eleven patients were stage IIA (36.7%) and 19 patients were stage IIIA (63.3%), meaning that all 30 patients showed postoperative pathologic findings of lymph node metastasis; 11 patients for N1 (36.7%) and 19 patients for N2 (63.3%). Only four patients (13.3%) had lymph node ratio (LNR) <0.1 between the metastatic and examined lymph nodes.

Romo1 expression and clinicopathologic parameters

Romo1 staining was typically observed in the cytoplasm of NSCLC cells. Romo1 expression levels were not normally distributed, and a representative example is shown in Figure 1 (average 7.6, range 0–15).

The process for determining the Romo1 cutoff level is described hereafter, and the cutoff was determined to be 8. With this cutoff, we classified 13 patients as being Romo1 low and 17 patients as Romo1 high. We evaluated the clinical parameters in these two groups, and NLR, CRP, CRP–albumin ratio, lymphatic invasion, LNR, and lymph node (N) stage were shown to be significantly associated with Romo1; *P*-values were 0.013, 0.04, 0.001, 0.024, 0.014, and 0.023, respectively (Table 1). These results were supported by the Mann–Whitney *U* test (Table S1).

Survival analyses

In the analyses described above, upregulated Romo1 was shown to be associated with high LNR, lymphatic invasion, and N2 stage rather than N1 stage, suggesting that Romo1 induces a high lymphatic metastatic tendency in patients with NSCLC. In addition, Romo1 was related to serologic biomarkers consistent with persistent inflammation which is believed to play an essential role in carcinogenesis and cancer progression. These findings suggest that Romo1 might augur an unfavorable prognosis for NSCLC patients. We evaluated the prognostic potential of Romo1, and several inflammatory markers in our cohort. The median follow-up time from curative resection was 38.8 months (range 1.4–115 months) and the median follow-up time among surviving patients was 50.6 months (range 5.4–115 months). Four patients out

Table 1 The proportions of patients with low and high Romo I expression according to clinicopathologic parameters

Parameters	Romo I		P-value
	<8 Patients, n (%)	≥8 Patients, n (%)	
Age (years)			0.31
<65	7 (23.3)	6 (20.0)	
≥65	6 (20.0)	11 (36.7)	
Sex			0.222
Male	11 (36.7)	11 (36.7)	
Female	2 (6.7)	6 (20.0)	
Smoking			0.367
Never	9 (30.0)	9 (30.0)	
Ever	4 (13.3)	8 (26.7)	
Pathology			0.352
Adenocarcinoma	8 (26.7)	11 (36.7)	
Squamous cell carcinoma	5 (16.7)	4 (13.3)	
Pleomorphic carcinoma	0 (0.0)	2 (6.7)	
Site			0.068
Right lung	2 (6.7)	8 (26.7)	
Left lung	11 (36.7)	9 (30.0)	
WBC ^a			0.374
<10,000/ μ L	13 (43.3)	16 (53.3)	
≥10,000/ μ L	0 (0.0)	1 (3.3)	
PLT ^a			0.141
<300,000/ μ L	12 (40.0)	12 (40.0)	
≥300,000/ μ L	1 (3.3)	5 (16.7)	
NLR ^a			0.013
<2.1	8 (26.7)	3 (10.0)	
≥2.1	5 (16.7)	14 (46.7)	
PLR ^a			0.087
<101	5 (16.7)	2 (6.7)	
≥101	8 (26.7)	15 (50.0)	
CRP ^a			0.04
<10.5 mg/L	12 (40.0)	10 (33.3)	
≥10.5 mg/L	1 (3.3)	7 (23.3)	
Albumin ^a			0.141
<3.5 g/dL	1 (3.3)	5 (16.7)	
≥3.5 g/dL	12 (40.0)	12 (40.0)	
CRP–albumin ratio ^a			0.001
<0.6	10 (33.3)	3 (10.0)	
≥0.6	3 (10.0)	14 (46.7)	
Lymphatic invasion			0.024
No	10 (33.3)	6 (20.0)	
Yes	3 (10.0)	11 (36.7)	
Vascular invasion			0.177
No	10 (33.3)	9 (30.0)	
Yes	3 (10.0)	8 (26.7)	
Perineural invasion			0.936
No	9 (30.0)	12 (40.0)	
Yes	4 (13.3)	5 (16.7)	
LNR			0.014
<0.1	4 (13.3)	0 (0.0)	
≥0.1	9 (30.0)	17 (56.7)	
N stage			0.023
N1	8 (26.7)	3 (10.0)	
N2	5 (16.7)	14 (46.7)	
T stage			0.713
T1	7 (23.3)	6 (20.0)	
T2	8 (26.7)	9 (30.0)	

Note: ^aBefore surgical resection.

Abbreviations: CRP, C-reactive protein; LNR, lymph node ratio; N, lymph node; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; PLT, platelet; Romo I, reactive oxygen species modulator-1; T, tumor; WBC, white blood cell.

of 11 with stage IIA and 15 patients out of 19 with stage IIIA underwent vinorelbine plus cisplatin chemotherapy as postoperative adjuvant treatment. At the time of analysis, 14 patients had died (46.7%), and cancer had recurred in 20 patients (66.7%) after surgery. The median follow-up time from recurrence was 16.1 months (range 0.2–77.3 months). Eleven patients with recurrence showed local recurrence, 13 patients had distant metastasis, and four patients had both. Fifteen patients out of 20 who had had recurrent disease underwent palliative chemotherapy; the chemotherapy regimens included pemetrexed/cisplatin, gemcitabine/cisplatin, pemetrexed monotherapy, taxane monotherapy, etoposide/cisplatin, and gemcitabine/carboplatin. Ten patients had no recurring disease and were still alive.

The median values for Romo1, NLR, PLR, and CRP–albumin ratio were 8.0 (range 0–15), 2.2 (range 0.7–5.9), 130.3 (range 55.9–351.6), and 0.87 (range 0.05–12.7), respectively. To examine the prognostic value of these markers, we chose 5-year survival as the stratifying point for ROC analysis, and the ROC curves of Romo1 and CRP–albumin ratio showed statistical utility being drawn above the reference line (Figure 2). We used the cutoff value that made the sum of sensitivity and specificity the highest. The determined optimal cutoffs for Romo1, NLR, PLR, and CRP–albumin ratio were 8, 2.1, 101, and 0.6, with an area under the curve of 0.715, 0.635, 0.630, and 0.755, respectively.

Analyses for DFS with respect to clinical parameters are shown in Table S2. The mean DFS was 47.4 months (95% CI 30.6–64.2 months). By the log-rank test, Romo1 ≥8 ($P=0.031$), NLR ≥2.1 ($P=0.024$), CRP ≥10.5 mg/L ($P=0.007$), and CRP–albumin ratio ≥0.6 ($P=0.004$) were significantly associated with shorter DFS (Figure 3).

Analyses of OS with respect to clinical parameters are also summarized in Table S2. Mean OS was 70.0 months (95% CI 53.5–86.6 months). By the log-rank test, Romo1 ≥8 ($P=0.011$), NLR ≥2.1 ($P=0.044$), PLR ≥101 ($P=0.026$), and CRP–albumin ratio ≥0.6 ($P=0.006$) were significantly associated with shorter OS (Figure 4). Based on the number of inflammatory markers identified as risk factors affecting OS, NLR, PLR, and CRP–albumin ratio, patients were divided into three groups (group A=0; group B=1 or 2; group C=3 risk factors), and the OS for each group was compared (Figure 5). OS was significantly shorter in group C compared with group A ($P_c=0.045$) or B ($P_c=0.024$).

Relationship between Romo I and VEGF

Romo I regulates mitochondrial release of ROS, which induces angiogenesis/lymphangiogenesis via hypoxia-inducible

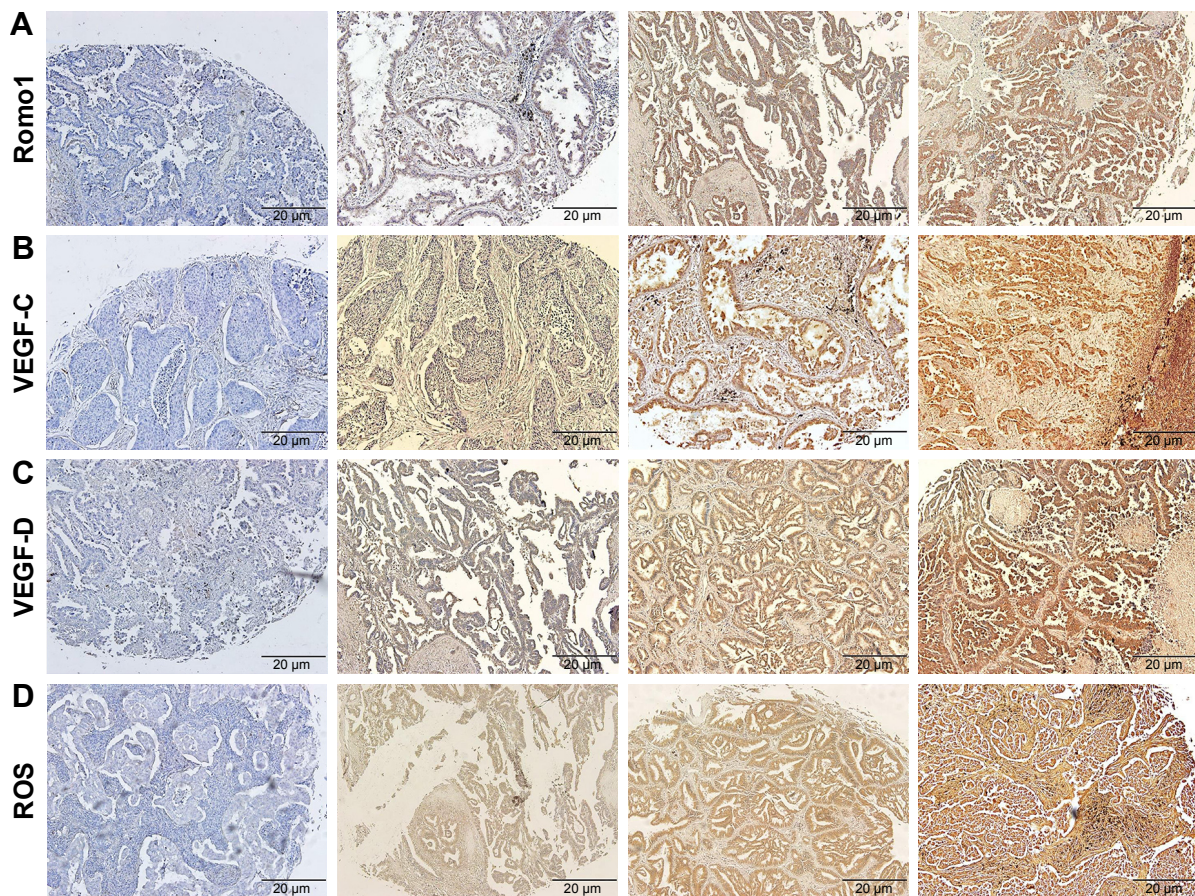


Figure 1 Representative IHC-stained tissue samples. Magnification, $\times 100$; **(A)** Romo1, negative control, IHC score of 2, 10, and 15; **(B)** VEGF-C, negative control, IHC score of 2, 9, and 15; **(C)** VEGF-D, negative control, IHC score of 2, 8, and 15; **(D)** ROS, negative control, IHC score of 1, 7, and 15.

Abbreviations: IHC, immunohistochemical; Romo1, reactive oxygen species modulator-1; ROS, reactive oxygen species; VEGF, vascular endothelial growth factor.

factor/VEGF signaling.¹⁵ Since Romo1 seems to influence a high lymphatic metastatic tendency, which is also associated with VEGF-C and VEGF-D, we investigated the correlation between Romo1 and VEGF-C, and VEGF-D in clinical samples by IHC staining (Figure 1). To determine whether a relationship exists between Romo1 expression, VEGF-C, VEGF-D, and ROS, correlation analyses were performed in paired tissue samples. Spearman's correlation analyses demonstrated significant positive correlations of Romo1 with VEGF-C ($P=0.008$, $R=0.478$) and ROS ($P=0.016$, $R=0.436$) (Figure 6).

Discussion

In this study, Romo1 overexpression was shown to be significantly correlated with early recurrence and an unfavorable prognosis in NSCLC patients who underwent curative surgery. There were also significant associations of unfavorable clinical outcomes with novel inflammatory biomarkers including NLR, PLR, and CRP–albumin ratio. Furthermore, there was significant association of Romo1 with high LNR, lymphatic invasion, and N2 stage rather than N1 stage,

demonstrating that the Romo1 expression level is related with lymphatic metastatic proclivity. In addition, Romo1 was associated with serum biomarkers of persistent inflammation, although this was likely predictable given that Romo1 controls oxidative stress. Considering both the oxidative stress/inflammation related with Romo1 and the lymphatic metastatic tendency associated with Romo1, we were able to speculate about a significant correlation between Romo1 and VEGF, and our subsequent analyses support it.

These results suggest that Romo1 might affect lymphatic metastasis of the primary tumor by regulating chronic inflammatory and oxidative stress (ROS)/VEGF signaling. Lymphatic metastasis related with Romo1 appears to be an essential correlate of the unfavorable clinical outcomes related to Romo1 overexpression. To our knowledge, this is the first study to provide a new hypothesis that lymphatic metastasis is the key reason for poor survival of NSCLC patients with elevated Romo1 and to propose persistent inflammation and oxidative stress/VEGF pathway as a possible mechanism by which Romo1 induces lymphatic metastasis of NSCLC.

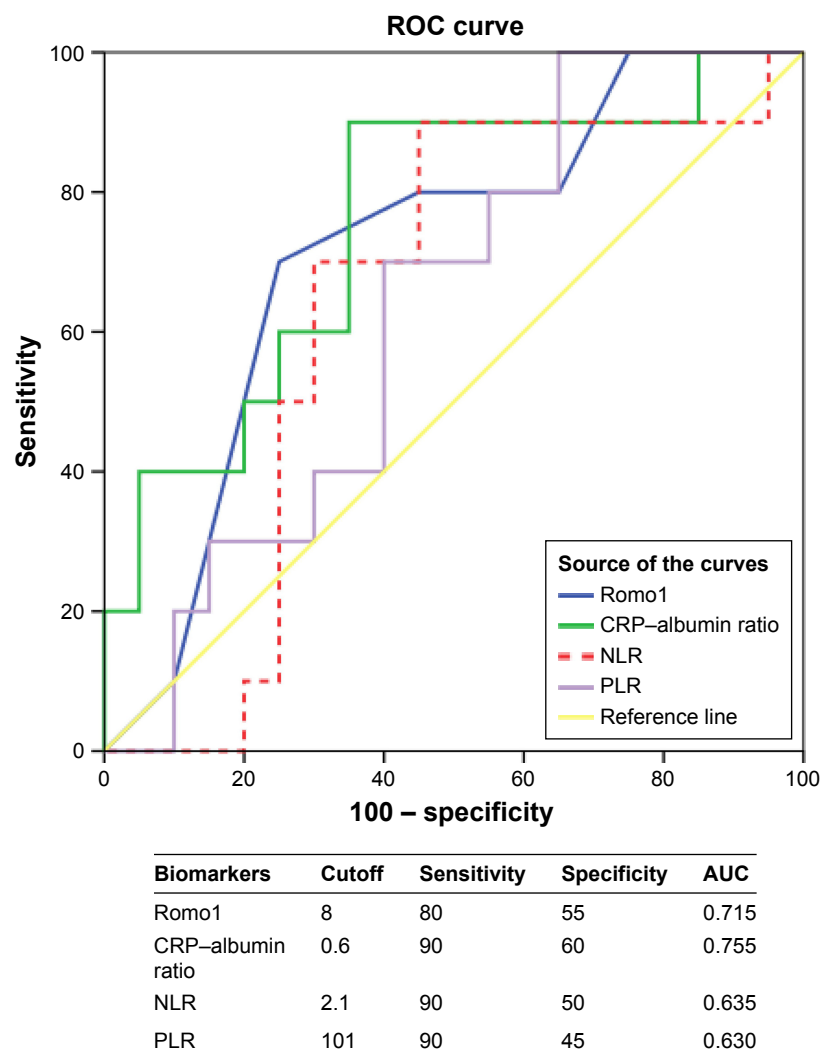


Figure 2 ROC analysis to set best cutoff for Romo1 and serologic inflammatory biomarkers (NLR, PLR, and CRP-albumin ratio).

Abbreviations: AUC, area under the curve; CRP, C-reactive protein; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; ROC, receiver operating curve; Romo1, reactive oxygen species modulator-1.

ROS levels were shown to be increased in various malignant diseases and associated with carcinogenesis and cancer progression;⁹ thus, studies on Romo1, a key regulator of mitochondrial ROS,¹⁰ have mainly been performed in the oncology field, especially in regard to Romo1's role in cancer cell invasiveness. In an earlier study, Romo1 knockdown induced downregulation of cell proliferation by blocking the extracellular signal-regulated kinases (ERK) pathway. Given that the ERK pathway is crucial for cell cycle entry, this finding suggests that Romo1 causes persistent oxidative stress via increasing ROS, resulting in carcinogenesis as well as cell proliferation.⁸ In another study, nuclear factor kappa B (NF- κ B) signaling pathway was downregulated by Romo1 knockdown, suggesting that Romo1 induces cancer cell invasion by upregulating the NF- κ B pathway. Increasing

invasiveness accelerated by Romo1 was interrupted by an inhibitor of κ B kinase (IKK), demonstrating the cooperative relationship between the NF- κ B pathway and Romo1.¹⁶

Since Romo1 was first found a decade ago, there have been studies demonstrating Romo1 as a prognostic biomarker in HCC, CRC, and NSCLC,^{11,13,14} but no study has investigated the mechanism by which Romo1 adversely affects prognosis of cancer patients. We have shown here that Romo1 expression is correlated with lymphatic invasion of the primary tumor, but not vascular or neural invasion; whether it is invasion of the primary tumor into lymphatics or vasculature currently depends on the lymphangiogenic or angiogenic vessels rather than the dormant vessels,¹⁷ and the progression to lymphangiogenesis or angiogenesis depends on specific and different pro-angiogenic factors.¹⁸

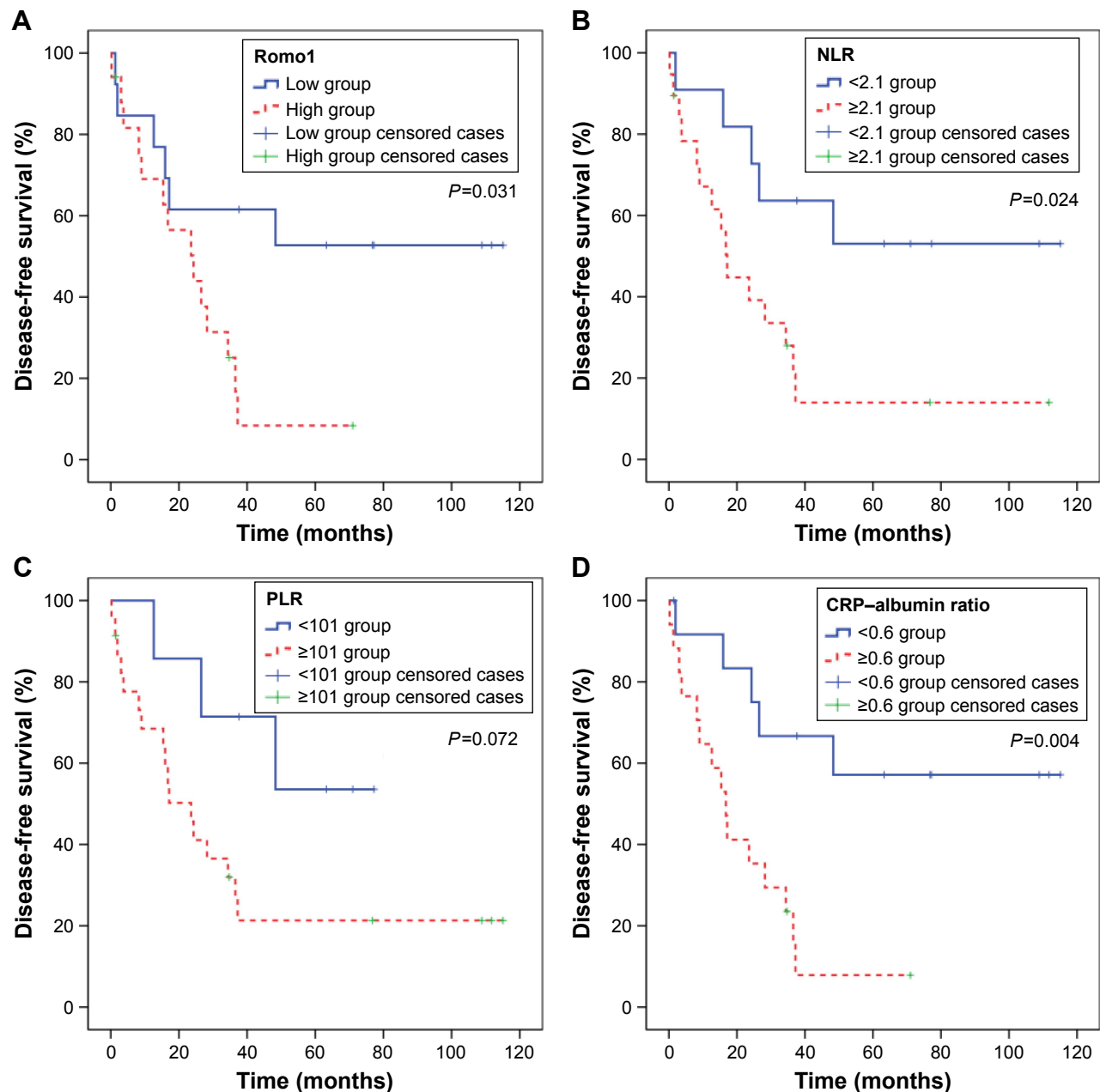


Figure 3 Kaplan–Meier curves of disease-free survival according to Romo1 (A), NLR (B), PLR (C), and CRP–albumin ratio (D).

Abbreviations: CRP, C-reactive protein; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; Romo1, reactive oxygen species modulator-1.

The VEGF gene family, which includes placental growth factor, VEGF-A, VEGF-B, VEGF-C, and VEGF-D,¹⁹ has signal proteins that induce angiogenesis and lymphangiogenesis. Among them, VEGF-C and VEGF-D are reported to promote the growth of tumor-associated lymphatic vessels and enhance lymphatic metastasis.²⁰ Based on both the fact that Romo1 is an upstream regulator of ROS, which has a cooperative relationship with the VEGF family,²¹ and the hypothesis that Romo1 may cause lymphangiogenesis, authors investigated the correlation of Romo1 with

VEGF-C and VEGF-D. In the current study, Romo1 was significantly related only with VEGF-C, suggesting that Romo1 might cause lymphatic metastasis via oxidative stress/VEGF-C signaling.

Romo1 could be a druggable target. Upregulated Romo1 has been reported to be associated with cancer invasiveness in HCC patients,¹³ and high lymphatic metastatic tendency in CRC patients¹⁴ as well as our study patients. Because the main driver of mortality in cancer is metastasis,²² targeting Romo1 might improve survival rates. In fact, considering

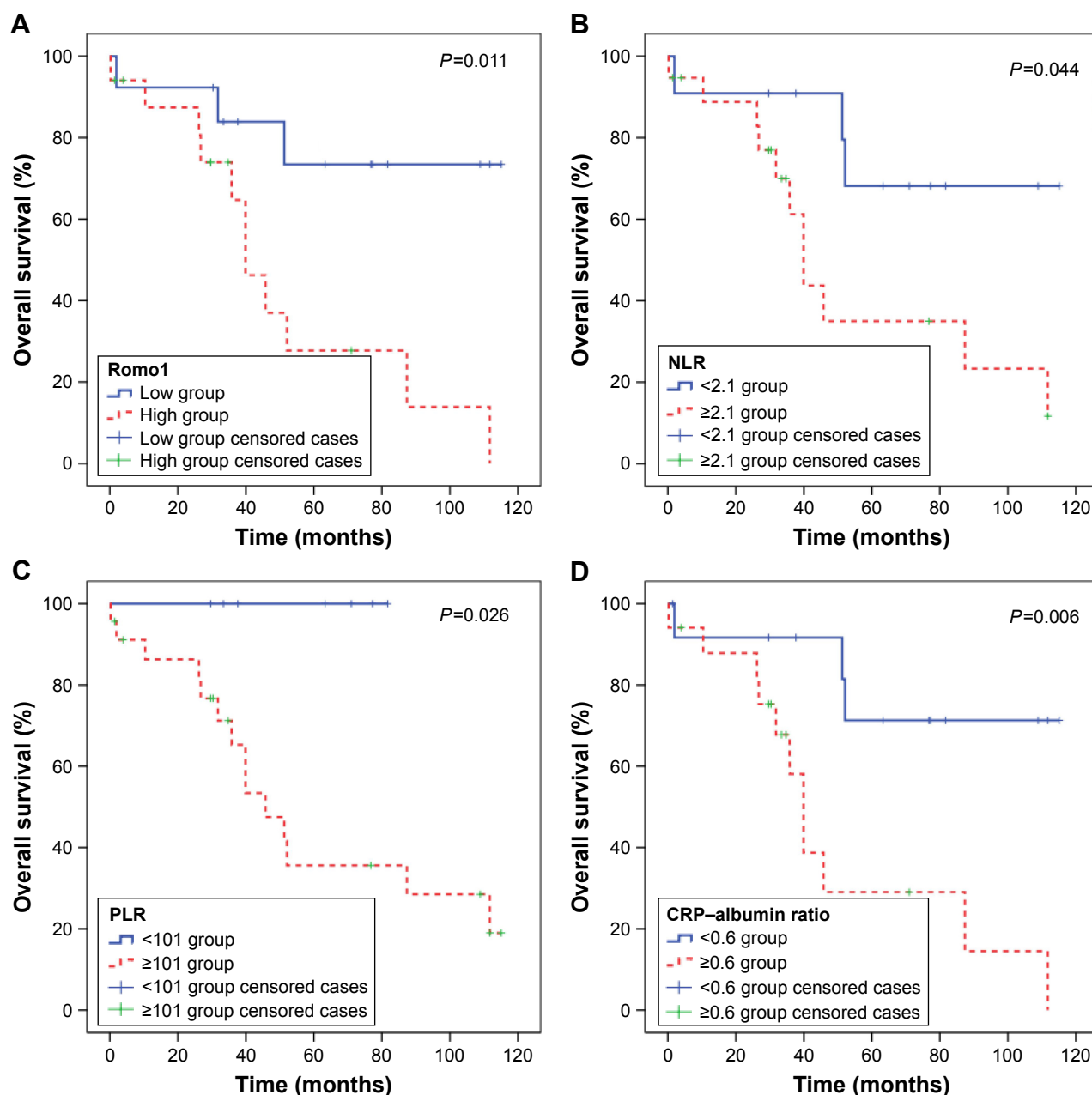


Figure 4 Kaplan-Meier curves of overall survival according to Romo1 (A), NLR (B), PLR (C), and CRP-albumin ratio (D).

Abbreviations: CRP, C-reactive protein; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; Romo1, reactive oxygen species modulator-1.

that chronic ROS stress is thought to be harmful in cancer progression, several antioxidants have been tested for their ability to prevent cancer invasion; however, all have shown disappointing results.²³ Rather than terminating ROS already released by mitochondria, regulating Romo1 may be a better option, making it a promising drug target.

In this study, Romo1 expression showed a significant correlation with novel inflammatory biomarkers, including NLR, PLR, and CRP-albumin ratio. Lately, these markers have been identified in several cancers as easily measurable

prognostic indicators, and many studies suggest that systemic persistent inflammation has an essential role in cancer biology.^{6,24} Considering that Romo1 is a regulator of ROS and correlates with inflammatory biomarkers, we think that carcinogenic inflammation could be regulated by targeting Romo1.

Inflammation has both tumor suppressing and tumor promoting features; acute inflammation inhibits cancer progression, while chronic inflammation induces carcinogenesis.⁷ This recognition guided clinical researchers into

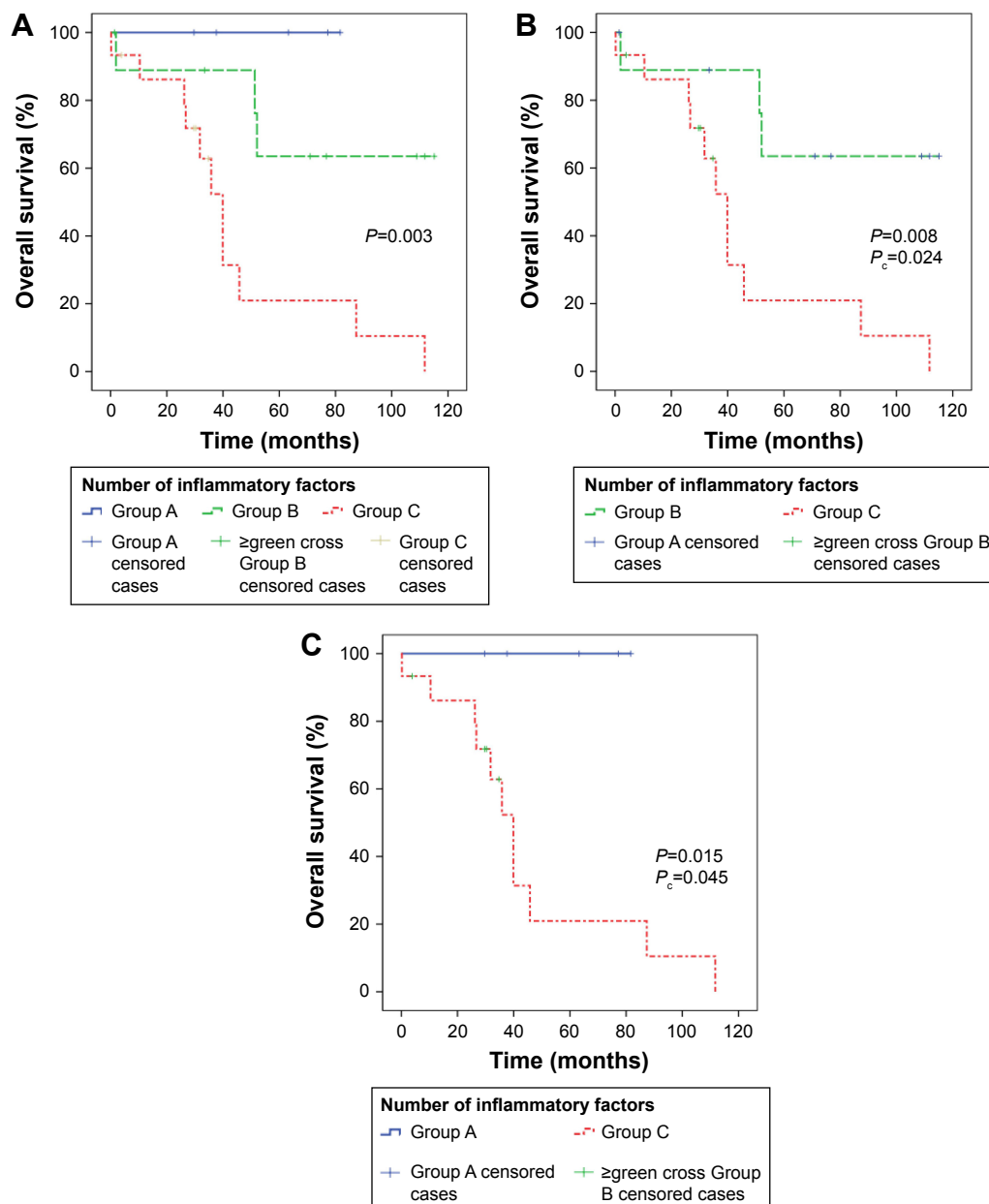


Figure 5 Kaplan-Meier curves of overall survival according to the number of inflammatory risk factors (A). Analyses using Bonferroni corrected P values (P_c) showed that overall survival was significantly shorter in group C compared with group B (B) or group A (C). Number of risk factors: group A =0; group B =1 or 2; group C =3.

the immuno-oncology field in pursuing research that promotes the cytotoxic properties of antigen-specific T-cells and/or suppresses cancer promoting inflammation.⁷ For over a decade, the immuno-oncology field has been in the spotlight, and admired for showing innovative practice changing successes.²⁵ However, given the absence of narrow patient selection based on a proper prediction of the therapeutic effect, immunotherapy has been considered as only stratified medicine rather than precision medicine like targeted agents. Microsatellite instability status and PD-L1 expression status have been studied as predictive markers in a few cancers including NSCLC and CRC.^{26,27} Romo1 also

has potential as a proper predictive biomarker for immune checkpoint inhibitors, considering both the fact that Romo1 induces chronic harmful inflammations, and the fact that the tumor microenvironment corrupted by chronic inflammations is one of the barriers of major anticancer effector cells in immuno-oncology, CD8⁺ T-cells.^{7,28} Based on these, Romo1 is expected to have possible roles as not only a predictive biomarker but also a target of drugs, which could be combined with immune checkpoint blockade therapy, because regulating harmful inflammation by targeting Romo1 could be beneficial for avoiding the resistance of immunotherapeutic approaches.

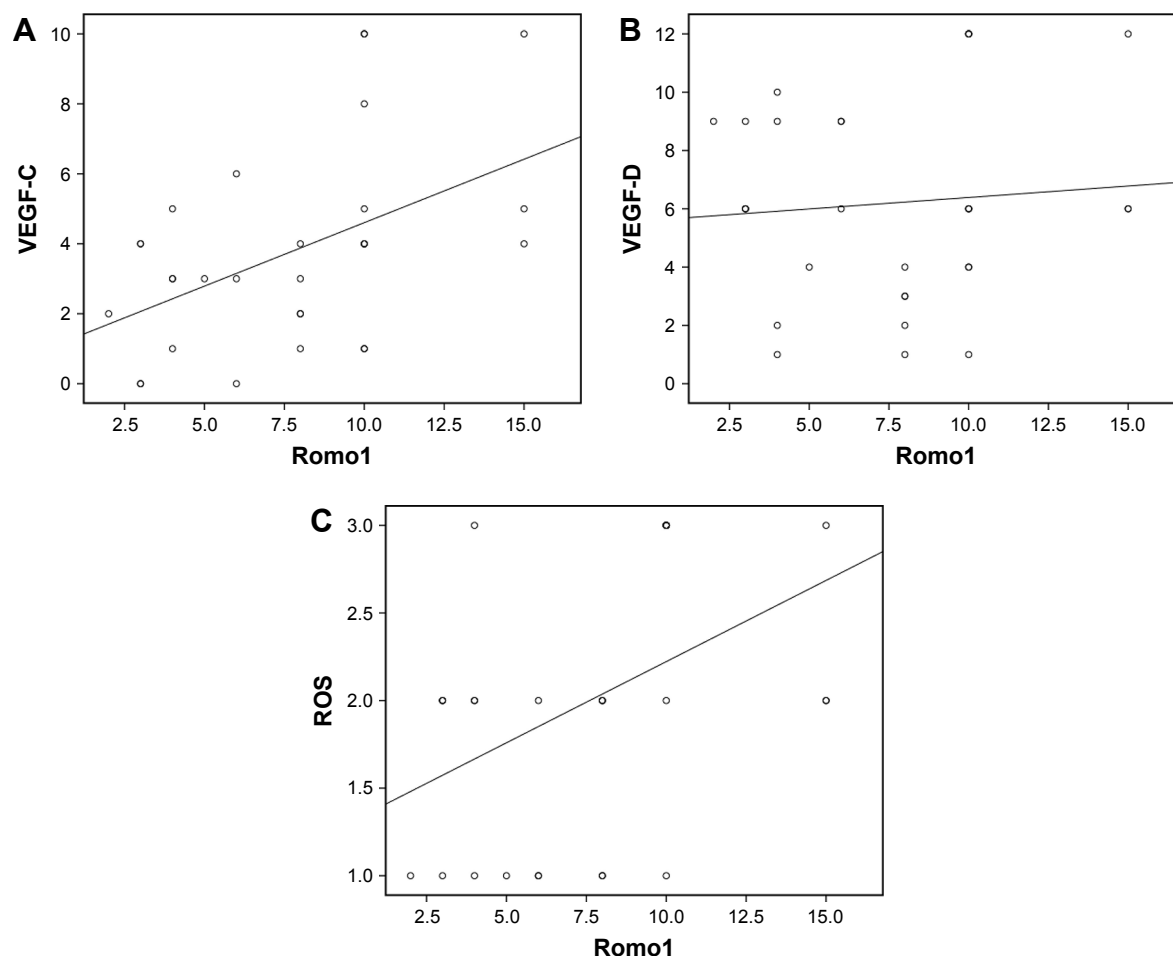


Figure 6 Spearman's correlation analyses evaluating correlation among Romo1, VEGF-C, VEGF-D, and ROS; Romo1 expression levels showed significant positive correlation with VEGF-C ($P=0.008$, $R=0.478$) (A) and ROS ($P=0.016$, $R=0.436$) (C) in tumor samples. (B) There was no significant correlation between Romo1 expression and VEGF-D levels.

Abbreviations: Romo1, reactive oxygen species modulator-1; ROS, reactive oxygen species; VEGF, vascular endothelial growth factor.

Limitations

This study has several limitations. First, IHC was used for the quantification of protein levels including Romo1 and VEGF, although the IHC method possibly has variant results. However, there is no standard quantification method established for the analysis of Romo1 expression and IHC is a generally conducted method for measuring protein expression. Not only that, protein expression analyses with a TMA could cause sampling error, because of a small portion of the whole tumor sample. Therefore, caution is needed not to overinterpret results of this study. Second, the results shown represent only associations rather than causations among serum inflammatory markers, overexpression of biomarkers including Romo1, VEGF, and ROS, and clinical variables related with lymphatic metastasis. In order to demonstrate the hypothesis of the mechanism by which Romo1 induces lymphatic metastasis of NSCLC by modulating persistent inflammation and oxidative stress (ROS)/VEGF signaling,

further properly designed in vitro studies are mandated. Third, the current study was conducted at a single center retrospectively with a relatively small sample size. Because our study cohort included patients with a relatively homogenous T stage, it is thought that possible bias in comparative analysis of Romo1 expression between the N1 group and N2 group (stage IIA and stage IIIA) was compromised. This might be the reason why the analyses were able to show statistical significance, even with a relatively small cohort. Our results should be validated by further prospective studies including many patients and using other quantification methods also.

Conclusion

The current study suggests a new hypothesis that Romo1 can induce lymphatic metastasis of NSCLC by modulating persistent inflammation/VEGF signaling. In this study, lymphatic metastasis related to Romo1 was shown to be a key reason

of unfavorable survival rates. With further research, antibodies targeting Romo1 could be designed and used as novel targeted agents. Romo1 could also be used as an indicator of harmful inflammatory conditions, so that researchers could expect possible barriers of immune checkpoint inhibitors and overcome them by targeting Romo1.

Disclosure

The authors report no conflicts of interest in this work.

References

- Jemal A, Center MM, DeSantis C, Ward EM. Global patterns of cancer incidence and mortality rates and trends. *Cancer Epidemiol Biomarkers Prev*. 2010;19(8):1893–1907.
- Jung KW, Won YJ, Kong HJ, et al. Cancer statistics in Korea: incidence, mortality, survival, and prevalence in 2012. *Cancer Res Treat*. 2015;47(2):127–141.
- Fauci AS, Longo DL, Kasper DL, Jameson JL, Loscalzo J, Hauser SL. *Harrison's Principles of Internal Medicine*. 17th ed. New York; London: McGraw-Hill Medical; 2008.
- DeSantis CE, Lin CC, Mariotto AB, et al. Cancer treatment and survivorship statistics, 2014. *CA Cancer J Clin*. 2014;64(4):252–271.
- Reck M, Rodríguez-Abreu D, Robinson AG, et al. Pembrolizumab versus chemotherapy for PD-L1-positive non-small-cell lung cancer. *N Engl J Med*. 2016;2016(375):1823–1833.
- Balkwill F, Charles KA, Mantovani A. Smoldering and polarized inflammation in the initiation and promotion of malignant disease. *Cancer Cell*. 2005;7(3):211–217.
- Coussens LM, Zitvogel L, Palucka AK. Neutralizing tumor-promoting chronic inflammation: a magic bullet? *Science*. 2013;339(6117):286–291.
- Na AR, Chung YM, Lee SB, Park SH, Lee MS, Do Yoo Y. A critical role for Romo1-derived ROS in cell proliferation. *Biochem Biophys Res Commun*. 2008;369(2):672–678.
- Wu WS. The signaling mechanism of ROS in tumor progression. *Cancer Metastasis Rev*. 2006;25(4):695–705.
- Chung YM, Kim JS, Yoo YD. A novel protein, Romo1, induces ROS production in the mitochondria. *Biochem Biophys Res Commun*. 2006;347(3):649–655.
- Lee SH, Choi SI, Lee JS, et al. Reactive oxygen species modulator 1 (Romo1) predicts poor outcomes in advanced non-small cell lung cancer patients treated with platinum-based chemotherapy. *Cancer Res Treat*. 2017;49(1):141–149.
- Lee SH, Min JW, Lee JS, et al. Reactive oxygen species modulator 1 (Romo1) overexpression is an independent predictor of poor survival in NSCLC patients who undergo surgical resection. *Lung Cancer*. 2015;87(1):45–52.
- Chung JS, Park S, Park SH, et al. Overexpression of Romo1 promotes production of reactive oxygen species and invasiveness of hepatic tumor cells. *Gastroenterology*. 2012;143(4):1084–1094.
- Kim HJ, Jo MJ, Kim BR, et al. Reactive oxygen species modulator-1 (Romo1) predicts unfavorable prognosis in colorectal cancer patients. *PLoS One*. 2017;12(5):e0176834.
- Kim YW, Byzova TV. Oxidative stress in angiogenesis and vascular disease. *Blood*. 2014;123(5):625–631.
- Lee S, Park YH, Chung JS, Yoo YD. Romo1 and the NF-κB pathway are involved in oxidative stress-induced tumor cell invasion. *Int J Oncol*. 2015;46(5):2021–2028.
- Dua R, Gui G, Isacke C. Endothelial adhesion molecules in breast cancer invasion into the vascular and lymphatic systems. *Eur J Surg Oncol*. 2005;31(8):824–832.
- Stacker SA, Achen MG, Jussila L, Baldwin ME, Alitalo K. Lymphangiogenesis and cancer metastasis. *Nat Rev Cancer*. 2002;2(8):573–583.
- Ferrara N, Gerber HP, LeCouter J. The biology of VEGF and its receptors. *Nat Med*. 2003;9(6):669–676.
- Karaman S, Detmar M. Mechanisms of lymphatic metastasis. *J Clin Invest*. 2014;124(3):922–928.
- Ushio-Fukai M. VEGF signaling through NADPH oxidase-derived ROS. *Antioxid Redox Signal*. 2007;9(6):731–739.
- Tam WL, Weinberg RA. The epigenetics of epithelial-mesenchymal plasticity in cancer. *Nat Med*. 2013;19(11):1438–1449.
- Ladas EJ, Jacobson JS, Kennedy DD, Teel K, Fleischauer A, Kelly KM. Antioxidants and cancer therapy: a systematic review. *J Clin Oncol*. 2004;22(3):517–528.
- Diem S, Schmid S, Krapf M, et al. Neutrophil-to-lymphocyte ratio (NLR) and platelet-to-lymphocyte ratio (PLR) as prognostic markers in patients with non-small cell lung cancer (NSCLC) treated with nivolumab. *Lung Cancer*. 2017;111:176–181.
- Burstein HJ, Krilov L, Aragon-Ching JB, et al. Clinical cancer advances 2017: annual report on progress against cancer from the American Society of Clinical Oncology. *J Clin Oncol*. 2017;35(12):1341–1367.
- Van Cutsem E, Cervantes A, Adam R, et al. ESMO consensus guidelines for the management of patients with metastatic colorectal cancer. *Ann Oncol*. 2016;27(8):1386–1422.
- Herbst RS, Baas P, Kim DW, et al. Pembrolizumab versus docetaxel for previously treated, PD-L1-positive, advanced non-small-cell lung cancer (KEYNOTE-010): a randomised controlled trial. *Lancet*. 2016;387(10027):1540–1550.
- Elinav E, Nowarski R, Thaiss CA, Hu B, Jin C, Flavell RA. Inflammation-induced cancer: crosstalk between tumours, immune cells and microorganisms. *Nat Rev Cancer*. 2013;13(11):759.

Supplementary materials

Table S1 Association between Romo I and clinicopathologic parameters

Parameters	Patients, n (%)	Average score of Romo I	P-value
All	30 (100)	8	
Age (years)			0.468
<65	13 (43.3)	7	
≥65	17 (56.7)	8	
Sex			0.251
Male	22 (73.3)	7	
Female	8 (26.7)	9	
Smoking			0.59
Never	18 (60.0)	8	
Ever	12 (40.0)	7	
Pathology			0.465
Adenocarcinoma	19 (63.3)	7	
Squamous cell carcinoma	9 (30.0)	7	
Pleomorphic carcinoma	2 (6.7)	12	
Site			0.169
Rt	10 (33.3)	9	
Lt	20 (66.7)	7	
Romo I ^a			N/A
<8	13 (43.3)	4	
≥8	17 (56.7)	10	
WBC ^a			0.953
<10,000/μL	29 (96.7)	8	
≥10,000/μL	1 (3.3)	8	
PLT ^a			0.205
<300,000/μL	24 (80.0)	7	
≥300,000/μL	6 (20.0)	9	
NLR ^a			0.013
<2.1	11 (36.7)	5	
≥2.1	19 (63.3)	9	
PLR ^a			0.022
<101	7 (23.3)	5	
≥101	23 (76.7)	8	
CRP ^a			0.03
<10.5 mg/L	22 (73.3)	7	
≥10.5 mg/L	8 (26.7)	10	
Albumin ^a			0.068
<3.5 g/dL	6 (20.0)	10	
≥3.5 g/dL	24 (80.0)	7	
CRP–albumin ratio ^a			0.002
<0.6	13 (43.3)	5	
≥0.6	17 (56.7)	9	
Lymphatic invasion			0.031
No	16 (53.3)	6	
Yes	14 (46.7)	9	
Vascular invasion			0.292
No	19 (63.3)	7	
Yes	11 (36.7)	8	
Perineural invasion			0.982
No	21 (70.0)	8	
Yes	9 (30.0)	8	
LNR			0.002
<0.1	4 (13.3)	3	
≥0.1	26 (86.7)	8	
N stage			0.057
N1	11 (36.7)	6	
N2	19 (63.3)	9	
T stage			0.866
T1	13 (43.3)	7	
T2	17 (56.7)	8	

Note: ^aBefore surgical resection.

Abbreviations: CRP, C-reactive protein; LNR, lymph node ratio; N, lymph node; N/A, not applicable; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; PLT, platelet; Romo I, reactive oxygen species modulator-1; T, tumor; WBC, white blood cell.

Table S2 Survival analyses for patients

Parameters	Patients, n (%)	Mean DFS (months)	P-value	Mean OS (months)	P-value
All	30 (100)	47.4		70	
Age (years)					0.079
<65	13 (43.3)	64.1	0.074	86.9	
≥65	17 (56.7)	32		56.7	
Sex					0.778
Male	22 (73.3)	49.4	0.778	71.3	
Female	8 (26.7)	31.6		67.8	
Smoking					0.396
Never	18 (60.0)	48	0.881	64.8	
Ever	12 (40.0)	46.6		75.5	
Pathology					0.221
Adenocarcinoma	19 (63.3)	44.1	0.325	69.1	
Squamous cell carcinoma	9 (30.0)	57.9		73.4	
Pleomorphic carcinoma	2 (6.7)	14.2		14.9	
Site					0.708
Rt	10 (33.3)	40	0.7	66	
Lt	20 (66.7)	51		72.2	
Romo I					0.011
<8	13 (43.3)	68.7	0.031	92.7	
≥8	17 (56.7)	24.2		51.6	
WBC ^a					0.657
<10,000/ μ L	29 (96.7)	48.1	0.797	N/A	
≥10,000/ μ L	1 (3.3)	28.2			
PLT ^a					0.392
<300,000/ μ L	24 (80.0)	50	0.644	73.8	
≥300,000/ μ L	6 (20.0)	31.7		57.8	
NLR ^a					0.044
<2.1	11 (36.7)	72.4	0.024	90.4	
≥2.1	19 (63.3)	30.5		56.4	
PLR ^a					0.026
<101	7 (23.3)	55.6	0.072	N/A	
≥101	23 (76.7)	37.7			
CRP ^a					0.146
<10.5 mg/L	22 (73.3)	59.6	0.007	76.9	
≥10.5 mg/L	8 (26.7)	16.4		56.2	
Albumin ^a					0.492
<3.5 g/dL	6 (20.0)	31.7	0.918	58.6	
≥3.5 g/dL	24 (80.0)	47.7		73	
CRP–albumin ratio ^a					0.006
<0.6	13 (43.3)	76.1	0.004	92.7	
≥0.6	17 (56.7)	21.6		50.9	
Lymphatic invasion					0.836
No	16 (53.3)	51.7	0.509	70.6	
Yes	14 (46.7)	41.3		69.4	
Vascular invasion					0.716
No	19 (63.3)	46.9	0.856	73.8	
Yes	11 (36.7)	48.3		60.2	
Perineural invasion					0.749
No	21 (70.0)	50.2	0.519	66.9	
Yes	9 (30.0)	38.8		74.9	
LNR					0.057
<0.1	4 (13.3)	86.9	0.104	N/A	
≥0.1	26 (86.7)	40.1			
N stage					0.299
N1	11 (36.7)	59.9	0.181	84.9	
N2	19 (63.3)	37.3		62.3	

Note: ^aBefore surgical resection.

Abbreviations: CRP, C-reactive protein; LNR, lymph node ratio; N, lymph node; N/A, not applicable; NLR, neutrophil to lymphocyte ratio; PLR, platelet to lymphocyte ratio; PLT, platelet; Romo I, reactive oxygen species modulator-1; WBC, white blood cell.

OncoTargets and Therapy**Dovepress****Publish your work in this journal**

OncoTargets and Therapy is an international, peer-reviewed, open access journal focusing on the pathological basis of all cancers, potential targets for therapy and treatment protocols employed to improve the management of cancer patients. The journal also focuses on the impact of management programs and new therapeutic agents and protocols on

patient perspectives such as quality of life, adherence and satisfaction. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Submit your manuscript here: <http://www.dovepress.com/oncotargets-and-therapy-journal>