



original reports

First-in-Human Phase I Study of MPO250, a First-in-Class DARPIn Drug Candidate Targeting VEGF and HGF, in Patients With Advanced Solid Tumors

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abstract

PURPOSE A first-in-human study was performed with MPO250, a DARPIn drug candidate. MPO250 specifically inhibits both vascular endothelial growth factor (VEGF) and hepatocyte growth factor (HGF) with the aim of disrupting the tumor microenvironment.

PATIENTS AND METHODS A multicenter, open-label, repeated-dose, phase I study was conducted to assess the safety, tolerability, and pharmacokinetics of MPO250 in 45 patients with advanced solid tumors. In the dose-escalation part, 24 patients received MPO250 as a 3-hour infusion once every 2 weeks at five different dose levels (0.5-12 mg/kg). Once the maximum tolerated dose (MTD) was established, 21 patients were treated with a 1-hour infusion (n = 13, 8 mg/kg, once every 2 weeks and n = 8, 12 mg/kg, once every 3 weeks) of MPO250 in the dose confirmation cohorts.

RESULTS In the dose-escalation cohort, patients treated with 12 mg/kg MPO250 once every 2 weeks experienced dose-limiting toxicities. Therefore, MTD was 8 mg/kg once every 2 weeks or 12 mg/kg once every 3 weeks. The most common adverse events (AEs) were hypertension (69%), proteinuria (51%), and diarrhea and nausea (both 36%); hypoalbuminemia was reported in 24% of patients. Most AEs were consistent with inhibition of the VEGF and HGF pathways. Exposure was dose-proportional and sustained throughout the dosing period for all patients (up to 15 months). The half-life was about 2 weeks. Signs of single-agent antitumor activity were observed: 1 unconfirmed partial response with a time to progression of 23 weeks and 24 patients with stable disease, with the longest duration of 72 weeks and a median duration of 18 weeks.

CONCLUSION MPO250 is a first-in-class DARPIn drug candidate with suitable tolerability and appropriate pharmacokinetic properties for further development in combination with other anticancer therapies.

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INTRODUCTION

We report here the first-in-human study of MPO250, a DARPIn drug candidate and novel biologic that specifically and simultaneously binds to and inhibits vascular endothelial growth factor (VEGF) and hepatocyte growth factor (HGF).¹

DARPIn molecules are a class of small, highly specific binding proteins that can be easily assembled in a multispecific format; this is especially attractive for oncology drug development where simultaneous targeting of several pathways could be advantageous. Several other DARPIn molecules are in the clinical and preclinical development stage.

The rationale for concomitant targeting of the VEGF- and HGF-driven signaling pathways is to produce

greater disruption of the tumor cell-supporting micro-environment and thus overcome the clinical shortcomings of mono-targeted VEGF and HGF inhibitors.²⁻⁹

MPO250 is a tri-specific molecule with individual domains binding VEGF and HGF with picomolar affinity and two domains binding human serum albumin (HSA) to increase its plasma half-life (Fig 1). Pre-clinical studies have shown MPO250 to have greater effects on tumor growth and angiogenesis than individual VEGF- and HGF-blocking DARPIn molecules and also demonstrated potentiation of the antitumor activities of cytotoxic and immunomodulatory agents.¹⁰ In addition, there is support from the literature for the potential to overcome treatment resistance, which is commonly caused by upregulation of the cMET pathway.^{11,12}

ASSOCIATED CONTENT

Data Supplement

Author affiliations and support information (if applicable) appear at the end of this article.

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CONTEXT

Key Objective

DARPin molecules are a class of small, highly specific binding proteins that can be easily assembled to bind multiple molecular targets. MP0250 is a tri-specific molecule with individual domains binding vascular endothelial growth factor (VEGF), hepatocyte growth factor (HGF), and human serum albumin to increase its plasma half-life. This first-in-human study of intravenously administered MP0250 sought to determine the maximum tolerated dose (MTD) and assess its dose-limiting toxicities, pharmacokinetics, safety, and tolerability.

Knowledge Generated

The MTD of MP0250 was 8 mg/kg once every 2 weeks or 12 mg/kg once every 3 weeks. Drug half-life was about 2 weeks. Most adverse events were consistent with inhibition of the VEGF and HGF pathways, most commonly hypertension, proteinuria, nausea, diarrhea, and hypoalbuminemia. Single-agent antitumor activity was observed, including 1 patient with an unconfirmed partial response.

Relevance

MP0250 is a first-in-class DARPin drug candidate with a tolerability profile and pharmacokinetic properties, making it suitable for further development in combination with other anticancer therapies.

This manuscript summarizes the phase I experience with MP0250 in patients with cancer and thus constitutes the basis of MP0250 clinical development as well as the development of other systemically administered DARPin molecules.

PATIENTS AND METHODS

Study Design and Treatment

This was a multicenter, open-label, phase I, dose-escalation, and expansion study of intravenously administered MP0250 (ClinicalTrials.gov identifier: [NCT02194426](https://clinicaltrials.gov/ct2/show/study/NCT02194426)). The primary objectives were to determine the maximum tolerated dose (MTD) and to assess dose-limiting toxicities (DLTs), pharmacokinetics (PKs), safety, and tolerability. The secondary objective was to characterize the immunogenicity of MP0250. Exploratory objectives included assessment of biomarkers and antitumor activity of MP0250.

The dose-escalation phase followed a traditional 3 + 3 study design including five dose cohorts (C1-C5) of 0.5, 1.5, 4, 8, and 12 mg/kg of MP0250 administered over a

3-hour infusion once every 2 weeks. The MTD was defined as the dose below the dose level that produces DLTs in $\geq 33\%$ of patients. Once the MTD was established, two dose expansion cohorts were opened to further characterize the safety and biological activity of the selected dose. In the expansion cohorts, MP0250 was administered as a 1-hour infusion once every 2 weeks (cohort 6, C6, 13 patients at 8 mg/kg) or once every 3 weeks (cohort 7, C7, 8 patients at 12 mg/kg). The switch from once every 2 weeks to once every 3 weeks was supported by emerging PK data, and the infusion duration was reduced from 3 hours to 1 hour for patient convenience and to align with future once-every-3-week combination studies. Patients who benefited from treatment could continue treatment until tumor progression, unacceptable toxicity, or withdrawal from this study.

A DLT was defined as any drug-related adverse event (AE) meeting DLT criteria (provided in the Data Supplement, online only) that occurred from the time of first dose until completion of the DLT period (defined as 1 week after the third infusion in the once every 2 weeks schedule and 1 week after the second infusion in the once every 3 weeks

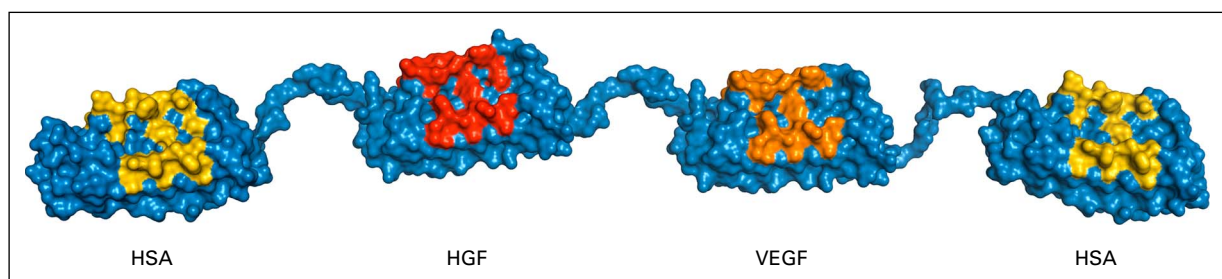


FIG 1. Structure of MP0250. HGF, hepatocyte growth factor; HSA, human serum albumin; VEGF, vascular endothelial growth factor.

schedule). A dose-escalation committee decided on dose escalation after all patients in a given cohort had completed the DLT period.

This study was approved by the ethics committees of participating institutions and appropriate regulatory authorities. All patients provided written informed consent. This study was conducted in compliance with the Declaration of Helsinki and guidelines for International Council for Harmonisation Good Clinical Practice.

Eligibility

Patients were at least 18 years old with advanced or metastatic solid tumor refractory to at least one line of prior standard treatment or for which no curative therapy was available. Patients had measurable or evaluable disease per RECIST version 1.1 and an Eastern Cooperative Oncology Group (ECOG) performance status of zero or one.

Safety Assessments

Routine clinical and laboratory assessments, including hematology, clinical chemistry, urinalysis, vital signs, and electrocardiography, were conducted at baseline and regularly throughout this study until up to 10 weeks after the last MPO250 administration.

AEs were continuously reported and coded using Medical Dictionary for Regulatory Activities version 17.0 terminology. AE intensity was graded according to the National Cancer Institute Common Terminology Criteria for Adverse Events (CTCAE) version 4.03.

PKs, Antidrug Antibodies, and Pharmacodynamics

Plasma samples for PKs, antidrug antibody (ADA), and cytokine assessment were collected at predefined time intervals. More detailed information is given in the Data Supplement.

Antitumor Activity

Radiologic tumor response assessments were performed by computed tomography or magnetic resonance imaging scans at screening and at 8-week intervals thereafter. Tumor responses were assessed using RECIST v1.1.

Statistical Analyses

This was an open-label phase I study. Sample sizes of at least 15 patients in the dose-escalation part (ie, at least three patients for the five doses planned) and 16 patients in the expansion part (eight per dose group) were considered sufficient to adequately address the objectives of this study. All patients exposed to MPO250 were included in the safety analysis, and all patients for whom at least one PK, pharmacodynamic (PD), or antitumor activity parameter could be reasonably assessed were evaluated for these parameters. All data were descriptively summarized.

Details of methods are provided in the clinical study protocol (Data Supplement).

RESULTS

Patient Characteristics

Twenty-four patients were enrolled in the dose-escalation phase, and 21 patients were enrolled in the expansion phase of this study. Patient characteristics of all 45 patients are summarized in Table 1. The most frequent tumor entities were colorectal cancer (31%) and ovarian cancer (13%). The ECOG performance status was 0 in 32 patients (71%) and 1 in 13 patients (29%). Twelve patients (27%) received previous anti-VEGF treatment. All patients completed at least one infusion of MPO250 (median, four infusions; range, 1-31 infusions).

MTD and DLT Evaluation

Treatment cohorts and DLTs are listed in Table 2. During the dose-escalation phase, four patients experienced DLTs. No DLTs were observed in the first two dose cohorts (0.5 and 1.5 mg/kg). In cohort 3 (4 mg/kg), one DLT occurred and this cohort was expanded to a total of six patients with no additional DLTs. In the next dose cohort (8 mg/kg), no DLT was reported in the first three treated patients. In cohort 5 (12 mg/kg), one of the three patients experienced two DLTs, leading to expansion of that cohort by two patients. After the second patient in this cohort experienced a DLT, the dose of the remaining patients was reduced to 8 mg/kg (two patients). Three more patients were enrolled at 8 mg/kg (cohort 4), of which one patient experienced

TABLE 1. Patient Demographics and Baseline Tumor Characteristics

Characteristic	All Patients (N = 45)
Median age, years (range)	63 (20-78)
Sex, n (%)	
Male	19 (42)
Female	26 (58)
ECOG performance status, n (%)	
0	32 (71)
1	13 (29)
Tumor type	
Colorectal cancer	14 (31)
Ovarian cancer	6 (13)
Breast cancer	3 (7)
Non-small-cell lung cancer (adenocarcinoma)	3 (7)
Head and neck cancer	3 (7)
Clear cell renal carcinoma	2 (4)
Spindle cell sarcoma	2 (4)
Others	9 (20)
Unknown primary	3 (7)

NOTE. Others: cervical, anal, mesothelioma, gallbladder, esophagus, melanoma, neuroectodermal, endometrial sarcoma, and endometrial carcinoma.

Abbreviation: ECOG, Eastern Cooperative Oncology Group

TABLE 2. Dose-Limiting Toxicities

DLTs, n (%)	CTCAE Grade	Cohort (Dose, mg/kg MP0250)								Total, N = 45
		C1 (0.5), n = 3	C2 (1.5), n = 3	C3 (4), n = 6	C4 (8), n = 7	C5 (12), n = 3	C4/5 (8/12), ^a n = 2	C6 (8), ^b n = 13	C7 (12) ^{b,c} n = 8	
Acute left ventricular failure	3	—	—	1 (17)	—	—	—	—	—	1 (2)
Blood creatinine increase	2	—	—	—	1 (14)	—	—	—	—	1 (2)
Blood urea increase	3	—	—	—	1 (14)	—	—	—	—	1 (2)
Gastrointestinal hemorrhage	3	—	—	—	—	1 (33)	—	—	—	1 (2)
Hypertension	3	—	—	—	—	1 (33)	—	—	—	1 (2)
Nephrotic syndrome	3	—	—	—	—	1 (33)	—	1 (8)	1 (13)	3 (7)
Thrombotic microangiopathy	4	—	—	—	1 (14)	—	—	—	—	1 (2)
Total number of patients ^d		—	—	1 (17)	1 (14)	2 (67)	—	1 (8)	1 (13)	6 (13)

NOTE. Unless indicated otherwise, infusions were administered over 3 hours every 2 weeks.

Abbreviations: C, cohort; CTCAE, Common Terminology Criteria for Adverse Events; DLTs, dose-limiting toxicities.

^aFirst infusion 12 mg/kg and subsequent infusions 8 mg/kg.

^bAdministration over 1 hour.

^cOnce-every-3-weeks dosing schedule.

^dOne patient may experience more than 1 DLT.

three DLTs. As no more than 17% of patients experienced a DLT at this dose level, the administration of 8 mg/kg once every 2 weeks (and its dose intensity equivalent of 12 mg/kg once every 3 weeks) was established as MTD for MP0250 monotherapy in oncology patients and consequently used in the expansion phase. In the dose expansion phase, two of the 21 patients experienced DLTs.

Safety

Treatment-emergent AEs that were reported in more than 10% of patients are shown in Table 3. The most frequent AEs were hypertension (69% of patients), proteinuria (51%), and diarrhea and nausea (both 36%). Hypoalbuminemia was reported in 24% of patients. Proteinuria was reported at a MP0250 dose \geq 8 mg/kg once every 2 weeks, while hypertension was observed at all tested dose levels. The majority of AEs were of grade 1 or 2. An infusion-related reaction was reported in one patient.

Grade 3 hypertension and proteinuria were reported in 16 and three patients (36% and 7%), respectively. The median onset was at day 21 for hypertension and day 62 for proteinuria. Median durations of the AEs were 8 days and 29 days for hypertension and proteinuria, respectively. No correlation between previous anti-VEGF treatment and the occurrence of these AEs was found. Treatment was discontinued due to proteinuria in six patients (13%), including one patient experiencing concomitant grade 2 hypertension.

Eleven patients (24%) experienced serious adverse reactions, as shown in the Data Supplement; those reported by more than one patient included nephrotic syndrome (four patients, 9%) and pulmonary embolism (three patients, 7%). Ten patients died during the treatment phase of this study, nine of them due to disease progression. The tenth patient died

from cardiac failure where MP0250 could not be completely excluded as being a contributing factor; however, this patient had several preexisting cardiac risk factors: significant exposure to anthracycline chemotherapy and prior mediastinal irradiation, hypertension, left bundle branch block, and dyslipidemia. Later, atrial fibrillation was also reported as an AE (CTCAE severity grade 2), and the patient died 65 days after the first and last doses of MP0250. This event was judged retrospectively to be possibly related to MP0250 administration.

Treatment was discontinued due to AEs in 14 patients (31%). The most frequent AEs leading to treatment discontinuation were nephrotic syndrome (four patients, 9%), malignant neoplasm progression (two patients, 4%), and proteinuria (two patients, 4%).

PKs and ADAs

PK parameters are shown in Table 4, and median concentration-time profiles of cohorts in the dose-escalation part are depicted in Figure 2. After MP0250 infusion, the maximum concentration (C_{max}) was followed by a decline of serum concentrations in a roughly monoexponential manner. The exposure-related parameters C_{max} and area under the curve (AUC) increased in proportion to the dose. The estimated values of clearance, volume of distribution, and half-life were similar between dose levels, suggesting a linear PK of MP0250 in the dose range from 0.5 to 12 mg/kg. The estimated ranges of the geometric means of clearance and volume of distribution in the steady state were 0.070-0.146 mL/h/kg and 27.5-59.9 mL/kg, respectively.

Following multiple-dose administration, MP0250 accumulated in the once-every-2-weeks and once-every-3-weeks dosing regimens. Steady-state conditions were established after approximately 6 weeks, corresponding to infusion 4

TABLE 3. Common Treatment-Emergent Adverse Events (in at Least 10% of Patients, Highest Grade Per Patient)

		Cohort (Dose, mg/kg MPO250)									
MedDRA System Organ Class	Preferred Term	CTCAE Grade	Cohort (Dose, mg/kg MPO250)								Total, N = 45
			C1 (0.5), n = 3	C2 (1.5), n = 3	C3 (4), n = 6	C4 (8), n = 7	C5 (12), n = 3	C4/5 (8/12), ^a n = 2	C6 (8), ^b n = 13	C7 (12) ^{b,c} n = 8	
GI disorders	Abdominal pain	All	—	—	1 (17)	2 (29)	1 (33)	—	2 (15)	1 (13)	7 (16)
	Constipation	All	—	—	1 (17)	—	1 (33)	—	2 (15)	2 (25)	6 (16)
	Diarrhea	All	1 (33)	2 (67)	2 (33)	3 (43)	1 (33)	1 (50)	3 (23)	3 (38)	16 (36)
		3	—	—	—	—	—	—	1 (8)	—	1 (2)
	Nausea	All	—	1 (33)	3 (50)	3 (43)	—	—	5 (39)	2 (25)	14 (31)
	Vomiting	All	1 (33)	—	2 (33)	4 (57)	—	—	3 (23)	2 (25)	12 (27)
General disorders	Asthenia	All	—	—	—	1 (14)	—	—	1 (8)	3 (38)	5 (11)
	Fatigue	All	1 (33)	2 (67)	1 (17)	4 (57)	2 (67)	1 (50)	2 (15)	1 (13)	14 (31)
		3	—	—	—	1 (14)	—	—	—	—	1 (2)
	Pyrexia	All	—	1 (33)	1 (17)	2 (29)	—	1 (50)	2 (15)	2 (25)	9 (20)
Laboratory abnormalities	Alanine aminotransferase increase	All	—	—	1 (17)	—	1 (33)	—	2 (15)	2 (25)	6 (13)
		3	—	—	1 (17)	—	—	—	—	—	1 (2)
	Blood CPK increase	All	—	—	—	1 (14)	—	1 (50)	3 (23)	1 (13)	6 (13)
		3	—	—	—	—	—	—	1 (8)	—	1 (2)
	Blood creatinine increase	All	—	—	2 (33)	2 (29)	—	—	3 (23)	—	7 (16)
	Platelet count decrease	All	—	—	1 (17)	4 (57)	—	—	3 (23)	—	8 (18)
	4	—	—	1 (17)	—	—	—	—	—	1 (2)	
Metabolism and nutrition	Decreased appetite	All	1 (33)	—	2 (33)	1 (14)	1 (33)	—	4 (31)	3 (38)	12 (27)
	Hypoalbuminemia	All	—	—	1 (17)	2 (29)	—	1 (50)	4 (31)	3 (38)	11 (24)
		3	—	—	—	—	—	—	1 (8)	—	1 (2)
Musculoskeletal	Back pain	All	1 (33)	—	1 (17)	1 (14)	1 (33)	—	2 (15)	2 (25)	8 (18)
Neoplasms	Malignant neoplasm progression	All	—	—	3 (50)	1 (14)	1 (33)	—	2 (15)	—	7 (16)
		3	—	—	1 (17)	—	—	—	—	—	5 (11)
		5	—	—	2 (33)	1 (14)	1 (33)	—	1 (8)	—	(2)
Nervous system disorders	Headache	All	—	—	1 (17)	1 (14)	1 (33)	1 (50)	2 (15)	3 (38)	9 (20)
Renal urinary disorders	Proteinuria	All	—	—	—	5 (71)	1 (33)	2 (100)	9 (69)	6 (75)	23 (51)
		3	—	—	—	—	1 (33)	1 (50)	1 (78)	—	3 (7)
Respiratory	Cough	All	1 (33)	1 (33)	1 (17)	3 (43)	—	1 (50)	3 (23)	1 (13)	11 (24)
	Dyspnea	All	—	1 (33)	1 (17)	2 (29)	1 (33)	—	1 (8)	—	6 (13)
		5	—	—	—	—	1 (33)	—	1 (8)	—	2 (4)
Vascular disorders	Hypertension	All	1 (33)	1 (33)	4 (67)	6 (86)	2 (67)	2 (100)	9 (69)	6 (75)	31 (69)
		3	—	1 (33)	3 (50)	3 (43)	1 (33)	1 (50)	5 (39)	2 (25)	16 (36)

NOTE. Unless indicated otherwise, infusions were administered over 3 hours every 2 weeks.

Abbreviations: C, cohort; CPK, creatine phosphokinase; CTCAE, Common Terminology Criteria for Adverse Events; DLTs, dose-limiting toxicities; n, number of cases; N, number of patients per cohort; MedDRA, Medical Dictionary for Regulatory Activities.

^aFirst infusion 12 mg/kg and subsequent infusions 8 mg/kg.

^bAdministration over 1 hour.

^cEvery 3-week dosing schedule.

TABLE 4. Pharmacokinetic Parameters of MP0250

		Cohort (Dose, mg/kg MP0250)													
		C1 (0.5)		C2 (1.5)		C3 (4)		C4 (8)		C5 (12)		C6 (8) ^a		C7 (12 mg/kg) ^{a,b}	
PK Parameter	Infusion Number	1	4	1	4	1	4	1	4	1	4	1	4	1	3
C_{max} ($\mu\text{g/mL}$)	n	3	3	3	2	6	5	7	3	5	n.d.	12	6	8	5
	Geometric mean	12.6	23.2	53.4	78.4	124	232	280	419	414	n.d.	246	481	379	498
	Coefficient of variation (%)	6.4	31	24	11	14	13	29	4.6	22		34	13	39	36
C_{min} ($\mu\text{g/mL}$)	n	3	3	3	1	5	4	6	3	5		10	6	7	4
	Geometric mean	3.97	8.53	20.6	29.8	46.6	119	101	188	102	n.d.	81.9	206	89.9	143
	Coefficient of variation (%)	13	10	9.9		29	31	37	16	25		28	20	47	47
t_{max} (h)	n	3	3	3	2	6	5	7	3	5		12	6	8	5
	Median	4.0	0.15	0.75	2.0	1.2	0.08	24	0.10	3.9	n.d.	3.9	4.1	2	2.3
	Minimum-maximum	2.2-8.0	0.03-2.3	0.42-4.5	0.08-4.0	0.12-4.1	0.08-4.0	0.07-71	0.03-3.8	2.0-120		0.03-71	0.18-166	0.07-166	0.13-8.0
AUC_{tau} ($\mu\text{g}^*\text{h/mL}$)	n	3	3	3	1	5	4	6	3	5		10	6	7	4
	Geometric mean	2,370	4,530	10,400	17,500	24,900	56,500	50,100	80,700	59,600	n.d.	48,800	114,000	81,900	130,000
	Coefficient of variation (%)	6.5	21	4.8		17	28	33	4.2	8.8		27	14	48	45
AUC_{inf} ($\mu\text{g}^*\text{h/mL}$)	n	3	3	3	1	6	3	4	2	4		9	5	4	4
	Geometric mean	3,760	7,610	18,900	27,200	42,800	99,400	82,300	154,000	87,300	n.d.	71,900	207,000	102,000	201,000
	Coefficient of variation (%)	13	12	10		22.8	76	29	23	15		27	26	21	46
CL (mL/h/kg)	n	3	3	3	1	6	4	4	3	4		9	6	4	4
	Geometric mean	0.146	0.120	0.0874	0.0945	0.0950	0.0707	0.097	0.100	0.137	n.d.	0.113	0.0703	0.118	0.0923
	Coefficient of variation (%)	13	23	10		20	28	31	4.3	15		27	14	21	45
V_{SS} (mL/kg)	n	3	3	3	1	5	2	4	2	4		8	5	4	3
	Geometric mean	49.7	42.8	36.0	29.5	35.9	53.7	40.9	40.9	41.0	n.d.	39.6	27.5	59.9	49.0
	Coefficient of variation (%)	3.4	40	9.7		19	47	43	28	10		24	33	22	73
AR – C_{max}	n	3		2		5		3				6		5	
	Mean	1.93		1.58		1.90		1.24		n.d.		1.64		1.43	
	Standard deviation	0.74		0.29		0.29		0.15				0.49		0.50	
AR – AUC	n	3		1		4		3				6		4	
	Mean	1.93		1.67		2.32		1.38		n.d.		2.08		1.74	
	Standard deviation	0.32				0.47		0.25				0.49		0.26	
$t_{1/2}$ (h)	n	3	3	3	1	6	3	4	2	4		9	5	4	4
	Median	252	260	293	225	250	244	281	276	215	n.d.	265	279	374	356
	Minimum-maximum	210-267	217-268	237-331		222-355	224-771	229-401	222-329	176-252		173-347	194-408	329-384	226-430

(continued on following page)

TABLE 4. Pharmacokinetic Parameters of MP0250 (continued)

PK Parameter	Infusion Number	Cohort (Dose, mg/kg MP0250)													
		C1 (0.5)		C2 (1.5)		C3 (4)		C4 (8)		C5 (12)		C6 (8) ^a		C7 (12 mg/kg) ^{a,b}	
		1	4	1	4	1	4	1	4	1	4	1	4	1	3
$t_{1/2}$ – effective (h)	n	3		1		4		3				6		4	
	Median	289		255		425		212		n.d.		388		389	
	Minimum-maximum	260-407				266-532		95-221				104-454		314-540	
$t_{1/2}$ follow-up (h)	n	2		3		3		5		2		8		2	
	Median	279		285		324		421		398		321		592	
	Minimum-maximum	255-303		214-340		284-405		369-628		311-486		177-2000		411-773	

NOTE. Unless indicated otherwise, infusions were administered over 3 hours every 2 weeks. Bold indicates the main values for each parameter.

Abbreviations: AR, accumulation ratio; AUC, area under the curve; AUC_{taur} , area under the curve for a dosing interval; AUC_{inf} , area under the curve from time 0 to infinity; C, cohort; CL, clearance; C_{max} / C_{min} , maximum/minimum concentration; n, number of patients for whom measurements were done; n.d., no data; $t_{1/2}$, half-life; t_{max} , time to maximum concentration; VSS, volume of distribution in the steady state.

^aAdministration over 1 hour.

^bOnce-every-3-weeks dosing schedule.

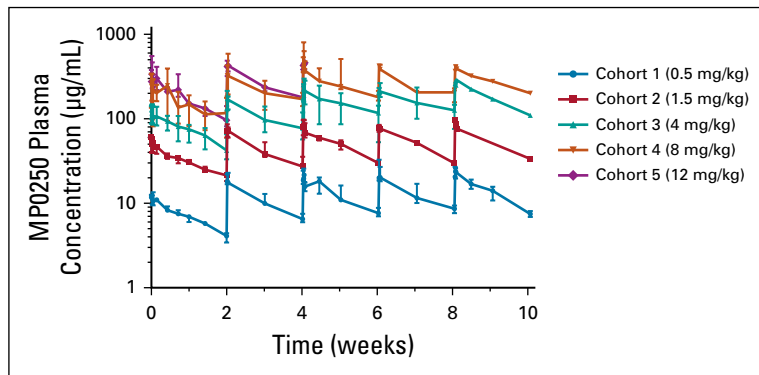


FIG 2. Pharmacokinetic traces of MP0250. Plasma concentration versus time profiles of cohorts 1-5 with once every 2 weeks dosing intervals. Numbers of patients per data point vary due to different numbers of patients in each cohort and drop out of patients after various periods ($n = 1-7$, median, max/min). A dose level of 12 mg/kg in cohort 5 was given to patients only for the first two cycles depicted in the graph.

in the once-every-2-weeks scheme and infusion 3 in the once-every-3-weeks scheme.

Half-life was calculated using three different approaches (in the follow-up period, based on accumulation,¹³ and during the dosing intervals). In all cohorts, a half-life of around 2 weeks was calculated during the follow-up periods (overall median of all patients $t_{1/2}$ follow-up 349 hours = 15 days) or based on the accumulation ratio (overall median of all patients $t_{1/2}$ effective 377 hours = 16 days). A similar half-life was calculated for cohort 7 over the dosing intervals of 3 weeks (overall median for cohort 7 was 374 hours = 16 days). Generally, a lower half-life was calculated during the shorter once-every-2-weeks dosing intervals for cohort 1 through cohort 6 (overall median for cohort 1 to cohort 6 and combined dose 1 and dose 4 was 252 hours = 11 days).

In general, no differences in PK results between males and females were observed.

ADAs were detected in 20 of the 42 evaluable patients and had no impact on the PK of MP0250. The occurrence of ADAs was independent of the MP0250 dose. In 2 of the 20 ADA-positive patients, the titer ratio (titer sample post dose/titer sample predose 1) was higher than 16. In one patient of cohort 2 (1.5 mg/kg, once every 2 weeks, 3-hour infusion), the titer ratio reached 256 (the highest observed value). This may be related to a deviation from the study protocol in which this patient did not have a filter in the infusion line during the first MP0250 infusion. One patient of cohort 6 (8 mg/kg, once every 2 weeks, 1-hour infusion) showed a maximal titer ratio of 64 although this decreased at a later time point. There were no apparent side effects associated with the presence of ADAs.

Pharmacodynamics

Binding of MP0250 to both its targets (VEGF-A and HGF) in plasma was demonstrated. For VEGF, this was shown by a reduction of plasma levels of free VEGF-A to undetectable for the duration of treatment (Data Supplement), and for HGF, it

was shown by total plasma HGF levels increasing in a dose- and time-dependent manner (Data Supplement). The increase in total HGF corresponded to an increase in MP0250-HGF complex in the circulation. Although the sample number was small, it appeared that MP0250-HGF complex formation might be plateauing around the dose of 4 mg/kg once every 2 weeks MP0250, thus possibly constituting the required dose for a maximum effect of MP0250 on the HGF biology.

PIGF (placental growth factor) levels decreased transiently in plasma after infusion of MP0250. There were no treatment-related changes observed for plasma levels of VEGF-C, angiopoietin-2, endoglin, fibroblast growth factor 2, interleukin (IL)-6, IL-8 and IL-10, tumor necrosis factor α , soluble VEGFR-2, or matrix metalloprotease 2 levels (data not shown).

Antitumor Activity

Among 40 patients evaluable for response, one patient (3%) had an unconfirmed partial response (uPR), 24 patients (60%) had stable disease (SD), and 15 patients (38%) had progressive disease.

The patient who had uPR had an anal cancer with lung metastases. The patient had received two prior lines of chemotherapy before entering into the trial. He received one infusion of 12 mg/kg followed by an infusion of 8 mg/kg once every 2 weeks and had PR at week 7 (30% reduction in target lesions), followed by SD at a subsequent tumor assessment with a time to progression of 23 weeks.

The median duration of SD was 18 weeks, with a maximum of 72 weeks in one patient with salivary gland carcinoma, who received MP0250 at the dose of 4 mg/kg (once every 2 weeks, 3-hour infusion).

The Kaplan-Meier analysis of progression-free survival (PFS) resulted in an overall median PFS of 15.1 weeks.

Of the 39 patients with a postdose assessment of the sum of longest diameters, 14 patients (36%) achieved some degree of tumor shrinkage (Data Supplement).

DISCUSSION

This study is the first clinical trial to evaluate the systemic administration of a new class of drugs, DARPin molecules, in patients with cancer. DARPin molecules are derived from protein binding domains found in ankyrin repeat proteins, a class of proteins widely used in nature for activities ranging from simple anchoring of proteins to enzyme inhibition.¹⁴ Binding domains with very high specificity and affinity can be selected and used as building blocks to create novel DARPin biologics that can combine multiple mechanisms of action in one molecule. MPO250 is a tri-specific molecule with four binding domains: one each for VEGF and HGF and two for HSA; albumin binding results in an extended plasma half-life, and the specific binding and neutralization of VEGF and HGF are responsible for therapeutic activity. The rationale to inhibit the VEGF and HGF pathways simultaneously is based on preclinical data that have shown that cMET can be upregulated to overcome resistance to VEGF inhibition and that simultaneous inhibition of these two pathways can overcome treatment resistance.^{9,11,12}

The predominant side effects of MPO250 observed in this first-in-human study were hypertension and proteinuria, both side effects that are characteristic of VEGF targeting drugs.¹⁵⁻¹⁷ Both side effects were dose-related, which is suggestive of an increasing inhibition of VEGF signaling with increasing doses. Although grade 3 hypertension was reported in about one-third of the patients, this AE was generally manageable and not treatment limited. Severe proteinuria was reported at the highest tested dose and was dose limiting in three patients (7%), and proteinuria appeared on average after 2 months of therapy. This AE warrants close monitoring in future clinical trials of MPO250, and since dose interruption is required in some cases, this should be reflected in the trial design. The dose of MPO250 to be used in combination studies could well be lower than the single-agent recommended phase II dose, particularly with other therapies that may have renal toxicities. With respect to biochemical signs of HGF inhibition, hypoalbuminemia, a class effect of HGF inhibitors, was frequently observed.⁵ PD data also demonstrated that both components of the DARPin molecule were active, resulting in a reduction of circulating VEGF below the limit of detection as well as a dose-proportional increase in HGF-MPO250 complexes. Thus, several pieces of data support the hypothesis that both the VEGF targeting and the HGF targeting components of MPO250 maintain their ability to bind and inhibit their specific targets when administered to patients, and the dose range tested here is highly relevant for the MPO250 pharmacology. With the established recommended

doses for further studies, MPO250 can now be tested in combination with other drugs to establish the role of concomitant VEGF/HGF inhibition as a potential remedy for patients with adaptive resistance.

The PK of MPO250 showed a dose-proportional increase in exposure (C_{max} and AUC) and demonstrated a serum half-life of approximately 2 weeks. Clearance was low with a range similar to values reported for therapeutic antibodies.^{18,19} The half-life of 2 weeks is significantly longer than what would be expected for a molecule in the size of MPO250 (65 kDa) and indicates that the majority of the drug is bound as intended to serum albumin. Consequently, convenient dosing intervals of once every 2, 3, or even 4 weeks can be used, depending on the combination therapy desired. No alteration in exposure was observed in any patient.

Although conducted in a population of heavily pretreated patients, this study demonstrated signs of single-agent activity. One patient showed a uPR and over half of the patients demonstrated stabilization of disease with the longest duration of 72 weeks.

In summary, this is the first study that demonstrates that rationally designed DARPin molecules, a new class of agents built on naturally occurring ankyrin repeat proteins, can be safely administered to patients with cancer. The molecule, MPO250, which contains three different components addressing three different targets, showed clinical and PD evidence that all intended targets are bound and that a simultaneous inhibition of VEGF and HGF is achieved. Based on the results of the phase I study, MPO250 is currently in phase II clinical evaluation in combination with bortezomib and dexamethasone in patients with refractory multiple myeloma²⁰ (ClinicalTrials.gov identifier: [NCT03136653](https://clinicaltrials.gov/ct2/show/study/NCT03136653)). Two additional DARPin drug candidates, MPO274, which binds to two different epitopes of the HER2 receptor and is capable of inducing apoptosis in HER2-expressing cell models, and MPO310 (AMG 506), a molecule that is designed to deliver 4-1BB activation to the tumor micro-environment via binding to fibroblast activation protein, have entered clinical testing (ClinicalTrials.gov identifiers: [NCT03084926](https://clinicaltrials.gov/ct2/show/study/NCT03084926) and [NCT04049903](https://clinicaltrials.gov/ct2/show/study/NCT04049903), respectively).

In a first-in-human clinical trial, MPO250, a rationally designed DARPin molecule, was found to have a manageable safety profile and showed signs of clinical activity supporting further clinical development especially in combination therapies. Furthermore, the data provide evidence that the DARPin platform could be a basis for a new class of drugs in which specific properties can be combined in a single molecule to provide innovative mechanisms of action.

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST**First-in-Human Phase I Study of MPO250, a First-in-Class DARPin Drug Candidate Targeting VEGF and HGF, in Patients With Advanced Solid Tumors**

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