## Fig. S1 Physician Treatment Selection Assessment Survey

## PHYSICIAN TREATMENT SELECTION ASSESSMENT SURVEY

Site Number:	Provider Name:	Date of Visit://
8-digit Subject ID:	Subject Name:	DD MM YYYY
Prescribed Therapy:		
	RAPY SELECTED FOR THE PATIE up to two (2) secondary reasons for N	
Primary Reason (SELECT ONE)  Active surveillance, no evidence of disease following procedure  Active surveillance, disease present Poor prognosis – supportive care, without Hospice  Poor prognosis – Hospice enrollment Unable to afford treatment Patient declined treatment Local therapy (metastasectomy, etc) Other (SPECIFY):	Secondary Reason (OPTIONAL)  Active surveillance, no evidence of disease following procedure  Active surveillance, disease present Poor prognosis – supportive care, without Hospice  Poor prognosis – Hospice enrollment Unable to afford treatment Patient declined treatment Local therapy (metastasectomy, etc) Other (SPECIFY):	Secondary Reason (OPTIONAL)  Active surveillance, no evidence of disease following procedure  Active surveillance, disease present Poor prognosis – supportive care, without Hospice  Poor prognosis – Hospice enrollment Unable to afford treatment Patient declined treatment Local therapy (metastasectomy, etc) Other (SPECIFY):
Please select the primary reason and patient. For combination therapies, p Primary Reason (SELECT ONE) Patient Characteristics Age Performance status/frailty	ERAPY SELECTED FOR THE PAT up to two (2) secondary reasons why lease indicate reasons in relation to ti Secondary Reason (OPTIONAL) Patient Characteristics Age Performance status/frailty	you chose the mRCC agent for the he combination as a whole.  Secondary Reason (OPTIONAL) Patient Characteristics Age Performance status/frailty
<ul> <li>□ Prognostic factors (MSKCC, Heng risk)</li> <li>□ Comorbidities (SPECIFY):</li> </ul>	<ul> <li>□ Prognostic factors (MSKCC, Heng risk)</li> <li>□ Comorbidities (SPECIFY):</li> </ul>	<ul> <li>Prognostic factors (MSKCC, Heng risk</li> <li>Comorbidities (SPECIFY):</li> </ul>
Likelihood of Clinical Benefit	Likelihood of Clinical Benefit  Potential for Tumor regression	Likelihood of Clinical Benefit  Potential for Tumor regression
Potential for Tumor regression     Overall survival/progression- free survival     Patient quality of life     Other (SPECIFY):	Overall survival/progression- free survival Patient quality of life Other (SPECIFY):	Overall survival/progression- free survival Patient quality of life Other (SPECIFY):
Overall survival/progression- free survival     Patient quality of life	<ul> <li>Overall survival/progression- free survival</li> <li>Patient quality of life</li> </ul>	<ul> <li>Overall survival/progression- free survival</li> <li>Patient quality of life</li> </ul>

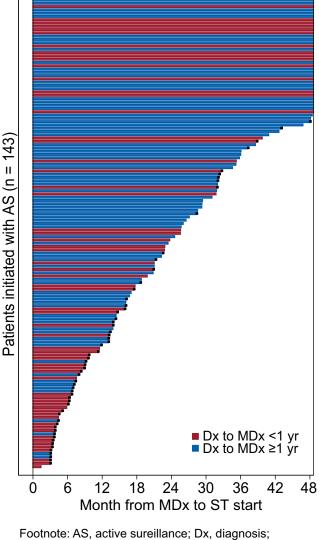
Prescribing Physician Signature: Physician Treatment Selection Assessment Survey Date:

(b)

## PHYSICIAN TREATMENT SELECTION ASSESSMENT SURVEY DISCONTINUING SYSTEMIC THERAPY

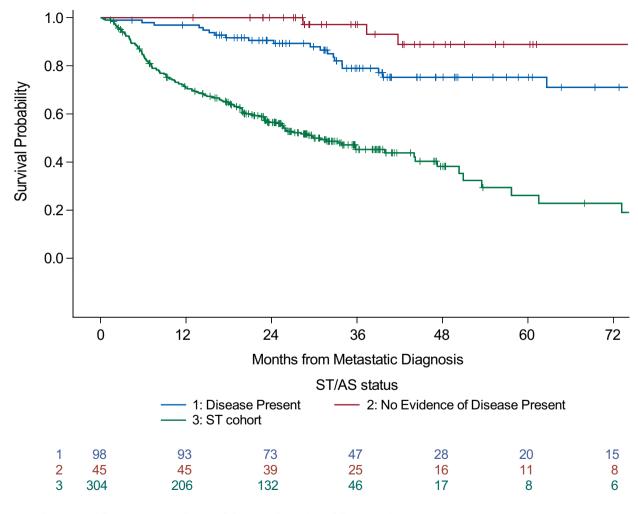
Site Number:	Provider Name:	Date of Visit:/ DD MM YYY
8-digit Subject ID:	Subject Name:	DD MM YYY
Prescribed Therapy:		
lease select the PRIMARY REASON	for discontinuing the most immediate prior	systemic therapy with the patient.
☐ Toxicity (GO TO C1)	☐ Disease Progression (GO TO C2	Other (GO TO C3)
	of SECONDARY toxicities and 1 ADDITIOn is sign to remove the patient from systematic	
Primary Toxicity (SELECT ONE)	Secondary Toxicity (SELECT ONE)	Secondary Toxicity (OPTIONAL)
Constitutional	Constitutional	Constitutional
☐ Fatigue/ Asthenia	☐ Fatigue/ Asthenia	☐ Fatigue/ Asthenia
☐ Arthralgia	☐ Arthralgia	☐ Arthralgia
Gastrointestinal	Gastrointestinal	Gastrointestinal
☐ Soreness in mouth/throat	□ Soreness in mouth/throat	☐ Soreness in mouth/throat
□ Diarrhea	☐ Diarrhea	☐ Diarrhea
☐ Nausea/vomiting	☐ Nausea/vomiting	☐ Nausea/vomiting
☐ Abdominal pain	☐ Abdominal pain	☐ Abdominal pain
Loss of appetite	□ Loss of appetite	□ Loss of appetite
Cardiovascular and Pulmonary	Cardiovascular and Pulmonary	Cardiovascular and Pulmonary
☐ Hypertension	☐ Hypertension	☐ Hypertension
☐ Cardiac dysfunction	☐ Cardiac dysfunction	☐ Cardiac dysfunction
Pneumonitis	☐ Pneumonitis	☐ Pneumonitis
Dermatologic/Skin	Dermatologic/Skin	Dermatologic/Skin
Soreness in hands/feet		☐ Soreness in hands/feet
☐ Rash	☐ Soreness in hands/feet ☐ Rash	☐ Rash
Hematologic and Laboratory	Hematologic and Laboratory	Hematologic and Laboratory
] Hemorrhage	☐ Hemorrhage	☐ Hemorrhage
☐ Anemia	☐ Anemia	□ Anemia
☐ Neutropenia	□ Neutropenia	☐ Neutropenia
☐ Thrombocytopenia	☐ Thrombocytopenia	☐ Thrombocytopenia
☐ Increased AST, ALT or Bilirubin		☐ Increased AST, ALT or Bilirubin
☐ Increased Creatinine	☐ Increased Creatinine	☐ Increased Creatinine
Other (SPECIFY):	□ None	Other (SPECIFY):
	Other (SPECIFY):	
treatment for the patient.	dicator(s) of DISEASE PROGRESSION	that led to discontinuation of system
New lesion(s) within already inv		
New lesion(s) in entirely new bo		
☐ Growth of existing lesion(s) (rac		
Symptomatic, disease related (	not treatment related toxicity)	
Other (SPECIFY):		
3. Please select the other reason	that led to discontinuation of systemic tr	eatment for the patient.
p Patient declined ongoing treatm		
Cost/ Unable to afford treatmen		
D Other (SPECIFY):	T/	
rescribing Physician Signature:		Date:

Fig. S2 The duration of time on AS



MDx, metastatic diagnosis; ST, systemic therapy. Each bar represents 1 patient. A square at the end of the bar indicates the patient initiated ST.

Fig. S3 Kaplan-Meier curves for OS in disease present, no evidence of disease present, and ST cohorts



Footnote: AS, active surveillance; OS, overall survival; ST, systemic therapy.