

# Clinical trials of multi target TKI for renal-cell carcinoma (advanced) in all type of patients

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## 1 TKI

Trial	Treatments	Patients	Trials design and methods
<b>apitolisib vs everolimus</b>			
Powles , 2014 n=NA	-	-	
<b>BNC105P + everolimus vs everolimus</b>			
Disruptor-1 n=NA	-	-	
<b>lenvatinib vs everolimus</b>			
Motzer , 2015 <i>ongoing</i> [NCT01136733] n=NA follow-up:	-	subjects with unresectable advanced or metastatic renal cell carcinoma following one prior VEGF-targeted treatment	
<b>sorafenib vs interferon alpha</b>			
Escudier , 2009 n=97/92 follow-up:	oral sorafenib 400 mg twice daily versus subcutaneous IFN-2a 9 million U three times weekly	patients with untreated, advanced renal cancer.	Parallel groups
<b>sunitinib vs interferon alpha</b>			
Motzer , 2007 [NCT00083889] n=375/375 follow-up:	repeated 6-week cycles of sunitinib (at a dose of 50 mg given orally once daily for 4 weeks, followed by 2 weeks without treatment) versus interferon alfa (at a dose of 9 MU given subcutaneously three times weekly).	patients with previously untreated, metastatic renal-cell carcinoma	
<b>pazopanib vs placebo</b>			
Sternberg , 2010 n=NA follow-up:	pazopanib versus placebo	treatment-naive and cytokine-pretreated patients with advanced renal cell carcinoma	Parallel groups double-blind
VEG105192 , 2010 [NCT00334282] n=290/145 follow-up:	-	treatment-naive and cytokine-pretreated patients with advanced renal cell carcinoma	
<b>sorafenib vs placebo</b>			

continued...

<b>Trial</b>	<b>Treatments</b>	<b>Patients</b>	<b>Trials design and methods</b>
<b>TARGET , 2007</b> [NCT00073307] n=451/452 follow-up:	continuous treatment with oral sorafenib (at a dose of 400 mg twice daily) versus placebo	patients with renal-cell carcinoma that was resistant to standard therapy	Parallel groups
<b>Ratain , 2006</b> n=NA follow-up:	-	patients with metastatic renal cell carcinoma	
<b>axitinib vs sorafenib</b>			
<b>AXIS (Rini) , 2011</b> [NCT00678392] n=NA follow-up:	-	second-line therapy in patients with metastatic renal cell cancer	
<b>Qin , 2012</b> n=NA	-	-	
<b>dovitinib vs sorafenib</b>			
<b>GOLD</b> [NCT01223027] n=284/286 follow-up:	dovitinib (500 mg orally according to a 5-days-on and 2-days-off schedule) versus sorafenib (400 mg orally twice daily)	patients with clear cell metastatic renal cell carcinoma who received one previous VEGF-targeted therapy and one previous mTOR inhibitor	open-label
<b>sunitinib vs sorafenib</b>			
<b>SWITCH</b> [NCT00732914] n=NA	-	-	
<b>tivozanib vs sorafenib</b>			
<b>TIVO-1 , 2013</b> [NCT01030783] n=260/257 follow-up:	tivozanib versus sorafenib	initial targeted therapy in patients with metastatic renal cell carcinoma	
<b>pazopanib vs sunitinib</b>			
<b>COMPARZ , 2013</b> [NCT00720941] n=557/553 follow-up:	continuous dose of pazopanib (800 mg once daily) versus sunitinib in 6-week cycles (50 mg once daily for 4 weeks, followed by 2 weeks without treatment)	patients with clear-cell, metastatic renal-cell carcinoma, first line	Parallel groups

## References

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**Disruptor-1, :**

**Motzer, 2015:**

**Escudier, 2009:**

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## 2 VEGFR, MET AXL TKI

Trial	Treatments	Patients	Trials design and methods
<b>cabozantinib vs everolimus</b>			
<b>METEOR , 2015</b> [NCT01865747] n=330/328 follow-up:	cabozantinib at a dose of 60 mg daily versus everolimus at a dose of 10 mg daily	patients with renal-cell carcinoma that had progressed after VEGFR-targeted therapy	Parallel groups open-label
<b>cabozantinib vs sunitinib</b>			
<b>CABOSUN , 2017</b> [NCT01835158] n=79/78 follow-up:	cabozantinib (60 mg once per day) versus sunitinib (50 mg once per day; 4 weeks on, 2 weeks off).	untreated clear cell mRCC and Eastern Cooperative Oncology Group performance status of 0 to 2 and were intermediate or poor risk per International Metastatic Renal Cell Carcinoma Database Consortium criteria	Parallel groups open-label

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## 3 About TrialResults-center.org

TrialResults-center is an innovative knowledge database that collects the results of RCTs and provides dynamic interactive systematic reviews and meta-analysis in the field of all major heart and vessels diseases.

The TrialResults-center database provides a unique view of the treatment efficacy based on all data provided directly from clinical trial results, offering a valuable alternative to personal bibliographic search, published meta-analysis, etc. Furthermore, it would allow comparing easily the various concurrent

therapeutic for the same clinical condition.

Rigorous meta-analysis method is used to populate TrialResults-center: widespread search of published and non published trials, study selection using pre-specified criteria, data extraction using standard form.

TrialResults-center is continually updated on a weekly basis. We continually search all new results (whatever their publication channel) and these news results are immediately added to the database with a maximum of 1 week.

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