

# Playing the Devil's Advocate: Should We Give a Second Chance to mTOR Inhibition in Renal Clear Cell Carcinoma? – ie Strategies to Revert Resistance to mTOR Inhibitors

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**Abstract:** In the last decade, the inhibition of the mechanistic target of Rapamycin (mTOR) in renal clear cell carcinoma (RCC) has disappointed the clinician's expectations. Many clinical trials highlighted the low efficacy and unmanageable safety profile of first-generation mTOR inhibitors (Rapalogs), thus limiting their use in the clinical practice only to those patients who already failed several therapy lines. In this review, we analyze the major resistance mechanisms that undermine the efficacy of this class of drugs. Moreover, we describe some of the possible strategies to overcome the mechanisms of resistance and their clinical experimentation, with particular focus on novel mTOR inhibitors and the combinations of mTOR inhibitors and other anti-cancer drugs.

**Keywords:** everolimus, temsirolimus, Rapa-Link, anti-angiogenics, autophagy

## Introduction

The treatment of metastatic Renal Cell Carcinoma (mRCC) has improved drastically over the last two decades. In fact, from a scarcity of specific therapeutic options, we assisted to the rise and the affirmation of a number of immune checkpoint inhibitors (ICIs) and small molecule antiangiogenic targeted agents, given either in combination or as a monotherapy, which dramatically changed the natural history of this once orphan disease.

Among all the molecular targets identified and successfully drugged in recent years, however, the mechanistic Target of Rapamycin (mTOR, previously mammalian Target of Rapamycin), has a pretty unique story. mTOR pathway controls several aspects of cell proliferation, migration, and metabolism, to an extent that is still largely unknown. The first identified mTOR inhibitor was, obviously, Rapamycin. This macrolide was first discovered in 1975 in a soil sample from the Easter Island of the South Pacific, also known as Rapa Nui (hence the name).<sup>1</sup>

In the early '90s, the mechanistic action of Rapamycin was clarified, leading to its application as an immunosuppressor and to the first in vitro studies on its anti-cancer properties. Hence, the first generation of Rapamycin derivatives, with reduced immunosuppressive effect, was developed. The two most promising molecules of this class, Temsirolimus and Everolimus (the so-called Rapalogs), have proven effective in several preclinical models. However, in clinical trials, they

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succeeded only in specific cancers, including mRCC. Moreover, the extent of this success was much less than expected. Rapalogs indeed showed a primarily cytostatic action, suggesting that their role should be disease stabilization rather than tumor regression. This has been attributed to the only partial inhibition of mTOR that fails to block the feedback loop between mTOR and its client kinase AKT, which arises as a mechanism of resistance.

In clinical trials involving mRCC, Rapalogs demonstrated superiority when compared with old therapeutic strategies, such as interferon-alpha. However, across the last decade, Rapalogs were heavily outclassed by multi-kinase inhibitors and immune checkpoint inhibitors. Therefore, they now find an application only after the failure of these drugs. On the other hand, more specific and potent mTOR inhibitors are currently being developed and tested, learning from the failures of the first generation. This review aims to analyze whether it is possible to overcome the resistance to mTOR inhibitors and what strategies should be preferred in the pursuit of this goal.

## The Role of mTOR in RCC

mTOR is an intracellular serine/threonine kinase that regulates several aspects of cell growth, proliferation, movement, and survival.<sup>2</sup> mTOR is able to react to many diverse stimuli (such as growth factors, cell nutritional status, and cellular stress) producing different responses. Specifically, mTOR exists in two distinct complexes, rapamycin-sensitive mTOR complex 1 (mTORC1) and rapamycin-sensitive mTOR complex 2 (mTORC2), that exert different effects, thus explaining the pleiotropy of this molecule. mTORC1 is formed by mTOR, regulatory-associated protein of mTOR (Raptor), G-protein  $\beta$ -subunit-like protein/LST8 (G $\beta$ L), and proline-rich AKT substrate 40 kDa (PRAS40); by modulating specific transcriptional factors, mTORC1 modulates lipids, protein and nucleotide biosynthesis, cell growth, autophagy, mitochondrial metabolism, and hypoxic response.<sup>3,4</sup>

On the other hand, mTORC2 is composed of G $\beta$ L, the protein rapamycin-insensitive companion of mTOR (Rictor), and mammalian stress-activated protein kinase (SAPK)-interacting protein (mSIN1); it plays a role in metabolic rewiring and cytoskeletal organization, being involved in cell movement.<sup>4</sup>

These intracellular serine/threonine kinases regulate many important cellular mechanisms<sup>5</sup> and the inability to regulate the mTOR pathway is emerging as a common feature in several malignancies. The abnormal activation

of the mTOR signaling pathway results in an unbalance in cell growth signals, leading to carcinogenesis. In most cancer cells, uncontrolled mTOR signaling provides a selective growth advantage.

The PIK3CA/AKT/mTOR pathway is one of the most commonly activated pathways in human cancers.<sup>5</sup> Mutations that trigger the pathologic activation of mTOR have been found in several types of cancer, albeit this axis is of uttermost importance in RCC,<sup>6</sup> a heterogeneous group of cancers originating from the nephron. There are several histotypes of kidney tumors, the most frequent being clear cell (65%), followed by the papillary subtype (20%) and the chromophobe subtype (5%).<sup>7</sup> Each histotype harbors its specific mutations, clinical characteristics, and therapeutic responses.<sup>8</sup>

Many studies aimed at identifying the gene mutations responsible for the development of RCC have been conducted over the years. A major role in clear cell RCC (ccRCC) genesis is played by the loss of function of the von Hippel-Lindau (VHL) gene, a tumor suppressor whose main role consists in inducing the degradation of the Hypoxia Inducible Factor (HIF). The second most studied mechanism involves the mutations affecting the PI3K/AKT/mTOR signaling pathway. These mutations either promote the overactivation of the PI3K/AKT/mTOR signaling cascade<sup>9</sup> or are responsible for the inhibition of the PTEN, a tumor suppressor gene that plays an inhibitory role on the aforementioned axis.<sup>10</sup> PTEN, PI3K, AKT, and mTOR represent fundamental checkpoints of the signal transduction pathway involved in the control of cell development and proliferation, being heavily implied in RCC pathogenesis and progression.<sup>11,12</sup> In fact, aberrations in the PI3K/AKT/mTOR pathway can be detected in 28% of clear cell RCC cases.<sup>13</sup> It's worth noting that mTOR mutation is particularly prevalent in highly vascularized tumor areas since vascularization allows nutrients and growth factors to activate the mTOR pathway.<sup>14-17</sup>

Mutations of the mTOR axis are also described in uncommon RCC subtypes, such as chromophobe RCC,<sup>18</sup> acquired cystic disease-associated RCC, eosinophilic solid and cystic RCC, RCC with fibromyomatous stroma and even in the low-grade oncocytic tumor.<sup>19</sup> Although the idea of a mTOR-driven RCC subclass is appealing, further research is needed in order to understand whether these mutations are to be considered driver or passenger mutations.

Whether these mutations can impact the survival of patients with RCC, is a debated topic. Fan et al

demonstrated that the increased expression of mTOR and its related proteins impacts positively on survival.<sup>20</sup> On the other hand, Ocana et al concluded that mutations of the mTOR axis, especially those implying the loss of PTEN, correlate with a significantly worse prognosis.<sup>21</sup> Regardless of its significance as a prognostic factor, mTOR remains a pivotal regulator of cell growth and proliferation, being thus considered a potential druggable target.

Indeed, in the past decades, different specific mTOR inhibitors were developed.

## mTOR Inhibitors in Current Clinical Practice

Since dysregulation of the PI3K/AKT/mTOR signaling pathway is frequently encountered in patients with RCC,<sup>22</sup> mTOR inhibitors were largely studied for the treatment of locally advanced or metastatic RCC.<sup>17,23</sup> Rapamycin was the first identified mTOR inhibitor. It acts by forming a complex with the protein FKBP12, which acts as the effective inhibitor of mTORC1.<sup>24</sup> From preclinic data, it's known that Rapamycin can slow cell growth and proliferation, or act synergistically with chemotherapy to induce cell death. Moreover, suppressing the mTOR pathway will slow down the consumption of external resources, inducing autophagy. Paradoxically, even an activator of the mTOR pathway could be therapeutically beneficial, because it would push cells to run out of energy and nutrients, and in the absence of stimuli, it could no longer maintain essential life processes.<sup>25,26</sup> In addition, Rapamycin was proven to inhibit metastasis of RCC in an in-vivo model.<sup>27</sup>

Although Rapamycin di per se was never extensively studied, as a therapeutic agent, in mRCC patients, its derivatives were largely used in this disease.

## Everolimus

Amato et al<sup>28</sup> first carried on a Phase II trial of Everolimus in patients with mcrRCC that received only 1 or no previous treatment lines. Everolimus reported an mPFS of 11.2 months, with an objective response rate (ORR) of 14% and a disease control rate (DCR) of 87%; this study represented the proof of principle of Everolimus activity in pre-treated mRCC patients.

In the RECORD-1 Phase III trial, Motzer et al<sup>29</sup> randomized mcrRCC patients progressing after one or more previous lines of therapy (including at least a previous

TKI, mostly Sunitinib) to receive either Everolimus or placebo. Everolimus yielded a significantly longer median PFS (4.9 months vs 1.9 months, HR 0.33,  $p < 0.00001$ ), with an acceptable safety profile. However, it should be noted that in this study, the crossover between arms was allowed, and was massive. As a consequence, no OS benefit was observed.

Everolimus was tested also in naive mcrRCC patients. In the RECORD-2 randomized phase II trial,<sup>30</sup> previously untreated mcrRCC patients were randomized to receive either the combination of Everolimus + Bevacizumab (an anti-VEGFR monoclonal antibody) or Interferon + Bevacizumab. The median PFS was 9.3 vs 10 months (HR = 0.90, 95% CI 0.71–1.15;  $P = 0.423$ ).

More recently, the RECORD-3 phase II trial<sup>31</sup> compared the sequence of Everolimus followed by Sunitinib with the reverse sequence (ie Sunitinib followed by Everolimus) in naive mRCC patients (including a subpopulation of non-clear cell RCC). Sunitinib upfront resulted in a higher median PFS (10 months vs 8 months) and ORR (27% vs 8%) when compared with Everolimus upfront. In addition, the combined median PFS of the two sequences was higher for that which started with Sunitinib (25.8 months vs 21.1 months). From all these findings it can be inferred that the place for Everolimus should be in the second-line, after the failure of a first-line with TKI. However, more recent trials furtherly scaled back the role of this drug.

In fact, the METEOR phase III trial<sup>32</sup> compared Cabozantinib and Everolimus in mcrRCC patients progressing after one or more VEGFR tyrosine kinase inhibitors. Cabozantinib reported a much better performance both in terms of mPFS (7.4 months vs 3.9 months) and ORR (17% vs 3%). In the same setting, Everolimus was compared with the immune checkpoint inhibitor Nivolumab.

Furthermore, the Checkmate-025 phase III trial<sup>33</sup> enrolled patients that received previous treatment with one or two regimens of antiangiogenic therapy and randomized them to receive either Nivolumab or Everolimus. Although the two drugs performed similarly in terms of median PFS (4.6 months vs 4.4 months), a significant difference was observed in the ORR (25% vs 5%) and median OS (25 months vs 19.6 months, HR = 0.73 (98.5% CI, 0.57 to 0.93;  $P = 0.002$ ), probably due to the development of long-lasting specific immune responses. Therefore, these trials suggest that in the common clinical practice, Everolimus should be administered to ccRCC

patients after the failure of immunotherapy and at least one line of TKI.

Everolimus was also tested in other RCC subtypes. The RAPTOR phase II trial evaluates the efficacy of Everolimus as a first-line treatment in a cohort of 92 patients with papillary RCC, reporting a median PFS of 4.1 months, which can be considered a clinical benefit since the treatment options for this variant are scarce.<sup>34</sup> Everolimus was also compared with Sunitinib in two major studies enrolling patients with papillary, chromophobe, or unclassified RCC: the ASPEN and the ESPN trials.<sup>35,36</sup> In both cases Sunitinib reported a better performance in terms of median PFS.

## Temsirolimus

Temsirolimus is the second most widely tested Rapalog. Hudes et al<sup>37</sup> tested this drug, within a randomized controlled phase III trial, in treatment-naïve mcrRCC patients having a poor prognosis (according to prespecified criteria), who were randomized to receive Temsirolimus, Interferon or a combination of the two agents. The Temsirolimus arm performed better than the others, with a median OS of 10.9 months versus 7.3 months in the Interferon arm and 8.4 months in the combination arm; this indeed was the very first (and for a long time, the only) randomized controlled trial which yielded an OS benefit in mcrRCC. On the other hand, the ORR was modest (8.6%).

In a subsequent trial,<sup>38</sup> Temsirolimus was compared with Sorafenib, in patients that failed the first-line therapy with Sunitinib. While median PFS did not statistically differ between the two groups (4.3 vs 3.9 months), there was an unexpected significant improvement in median OS in the Sorafenib arm (12.3 vs 16.6 months, HR, 1.31; 95% CI, 1.05 to 1.63; two-sided  $P = 0.01$ ).

Although Temsirolimus was not as widely tested as Everolimus, available data suggest that it could fit in a similar place in the clinical management of mcrRCC.

As for uncommon histologies, the CESAR trial compared Sunitinib and Temsirolimus in patients with papillary, chromophobe, medullary or unclassified RCC. Sunitinib outperformed Temsirolimus in terms of mPFS (13.6 vs 9.3 months, HR 1.75, 95% CI, 0.70–4.46).<sup>39</sup>

## Critical Appraisal of mTOR Inhibitors in mRCC

Upon a cursory glance, one of the weakest aspects of Rapalogs seems to be their relatively low ORR. As

a matter of fact, patients that experience shrinkage of lesions are less than 10% in all the aforementioned trials.

Interesting data derive from the analysis of those patients. Indeed, Voss et al<sup>40</sup> performed an elegant translational study on the tumors of patients that had long-lasting responses to Rapalogs (median duration: 28 months) and found genomic alterations with activating effect on mTOR signaling in the vast majority of them. Although it can fuel the idea that mTOR mutations could have a predictive role, this study has its limit in the low numerosity of the population and in its retrospective design. More recently, Roldan-Romero et al<sup>41</sup> performed genomic and immunohistochemical analysis on a wider cohort of patients that received Rapalogs. Their findings confirmed that mTOR pathway mutations and PTEN loss were significant predictors of partial responses. This data was confirmed also by Kwiatkowski et al<sup>42</sup> that studied a population of 79 mRCC patients treated with Rapalogs. Mutations in mTOR and its client network resulted to be more frequent among responders than non-responders. This evidence corroborates the idea that a strict population of Rapalogs responders exists among patients with mRCC, and that it could be selected by finding the correct biomarker. However, it should be noted that all these studies are flawed by their retrospective design.

The only known prospective clinical trial evaluating this aspect, was recently completed by Adib et al.<sup>43</sup> In this histology-agnostic trial, a pan-cancer cohort of patients with mTOR mutations was treated with Everolimus, obtaining a disappointing ORR of 7%. Therefore, even if mTOR mutations could have a role in the prediction of response to Everolimus, it is likely that they are not the only factor in the equation.

Nevertheless, even if mTOR mutations were confirmed to be the perfect biomarker, Rapalogs are still flawed by their safety profile. Common adverse events include hematological complications (thrombocytopenia, leukopenia, neutropenia, lymphopenia, and anemia), nausea, mucositis, diarrhea.<sup>44</sup> Beyond these adverse events, which are common across different anticancer agents, one of the most feared complications associated with the use of mTOR inhibitors is noninfectious pneumonitis, which can result in acute respiratory distress. In a randomized trial<sup>45</sup> comparing Everolimus vs placebo in patients with mcrRCC, 16% developed grade 2 and 3.6% grade 3 noninfectious pneumonitis in the Everolimus group.

In conclusion, in mRCC Rapalogs are drugs endowed with a relative and unpredictable efficacy, and an often

challenging safety profile. Given the premise of their extraordinary in-vitro results, their clinical performance was disappointing. The reasons behind this debacle have to be researched in the rapid development of resistance, a complex phenomenon, that arises from multiple fronts.

## Mechanisms of Resistance to mTOR Inhibitors

The main pathways and mechanisms implied in resistance to mTOR inhibitors are displayed in Figure 1.

### PI3K/AKT

As stated before, Rapalogs bind FKBP12, leading to the inactivation of pivotal proteins of the mTORC1 complex, while mTORC2 remains unaffected.<sup>46</sup> It has been demonstrated that mTORC1 inhibition leads to the reduction of Rictor (mTORC2 functional repressor) phosphorylation.<sup>47</sup> This will cause disinhibition of mTORC2, with an increase in its activity and a consequential hyperactivation of AKT.<sup>48</sup>

Moreover, the mTORC1 signal is able to activate, in a GSK3 (glycogen synthase kinase 3)-dependent way, the adapter protein GRB10, which mediates the PI3K-AKT negative feedback.<sup>49</sup> It follows that mTOR inhibition would lead to a loss of the PI3K-AKT negative feedback.

In addition, mTORC1 is indirectly responsible for the phosphorylation and subsequent degradation of IRS1 (insulin receptor substrate 1), which has an important role in facilitating insulin signaling,<sup>50</sup> which is implied in many anabolic mechanisms. It's therefore reasonable to hypothesize that the inhibition of mTORC1 caused by Rapalogs will disrupt the PI3K-AKT negative feedback and it will amplify the effects of insulin receptor signaling, thus leading to the activation of the MAP kinases and increasing cell proliferation, angiogenesis, survival, and a metabolic rewiring directed toward anaerobic glycolysis. This could be one of the reasons behind the poor performance of this drug class.

### Hypoxia Inducible Factor

HIF is strongly related to RCC tumorigenesis.<sup>51</sup> Under normal oxygen concentrations, the subunit HIF-1 $\alpha$  and HIF-2 $\alpha$  are constantly degraded by a VHL-dependent mechanism involving the proteasome.<sup>52</sup> In many cases, however, RCC is characterized by a loss of VHL,<sup>53</sup> resulting in persistent activation of HIF. In this scenario, HIF- $\alpha$  subunits will be able to enter the nucleus even if a normal concentration of oxygen exists, and it will bind HIF- $\beta$ , thus promoting the expression of genes implied in cell survival, apoptosis inhibition, and anaerobic metabolic rewiring.<sup>54</sup> However, it has been demonstrated that,

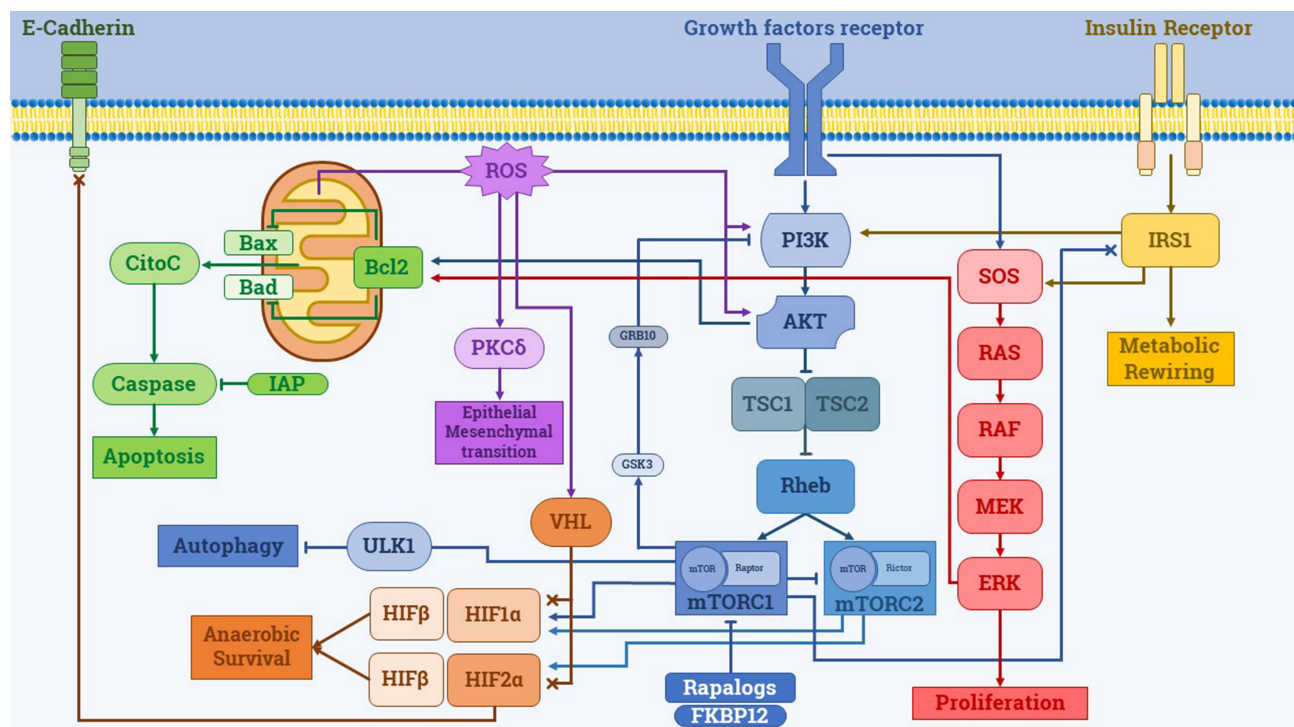


Figure 1 Main mechanisms involved in resistance to Rapalogs.

while HIF-1 $\alpha$  can exert both pro-proliferative and anti-proliferative effects, HIF-2 $\alpha$  has only pro-proliferative effects.<sup>55</sup> Moreover, HIF-2 $\alpha$  is always overexpressed in VHL-defective tumors, while HIF-1 $\alpha$  lacks in one-third of them.<sup>56</sup>

Interestingly, HIF- $\alpha$  subunits translation is stimulated by the PI3K/ATK/mTOR pathway.<sup>48</sup> In particular, while HIF-1 $\alpha$  expression depends on both mTORC1 and mTORC2, HIF-2 $\alpha$  depends only on mTORC2.<sup>57</sup> This data suggests that Rapalogs inhibition, which focuses on mTORC1, will not affect HIF-mediated RCC cells' survival improvement. Furthermore, it has been shown that E-cadherin is a target of the mTORC2/HIF-2 $\alpha$  axis. In particular, E-Cadherin is down-regulated by mTORC2/HIF-2 $\alpha$ , leading to the loss of contact inhibition and promoting cell migration and epithelial-to-mesenchymal transition.<sup>58</sup>

## Bcl2/IAP and Apoptosis Evasion

The antiapoptotic protein Bcl2 acts by regulating the permeability of the mitochondrial membrane and the cytoplasmatic release of cytochrome c. The role of Bcl2 in inhibiting apoptosis induced by Rapalogs is well-known and it has been extensively described.<sup>59</sup>

Another mechanism involves Survivin, a member of the IAP family, that acts by directly inhibiting caspases. There is indirect evidence that Survivin plays an important role in the evasion Rapalogs mediated apoptosis. It has been demonstrated that Rapalogs-mediated apoptosis can be greatly enhanced by reducing Survivin levels.<sup>60</sup>

## Mitogen-Activated Protein Kinases (MAPK) Pathway

MAP kinases are a family of intracellular kinases involved in proliferation, differentiation, mitosis, cell survival and apoptosis. Among MAPK c-Jun N-terminal kinases (JNKs) and p38 MAPKs (p38s) are usually triggered by cellular stresses and inflammatory cytokines signaling, while extracellular signal-regulated kinases (ERKs) are stimulated by growth factors.<sup>61</sup>

Inhibition of mTOR was demonstrated to induce an activation of the MAP-kinase ERK via a PI3K-dependent feedback loop. In particular, strong activation of the MAPK signaling was observed *in vitro*, after the administration of Everolimus. In the same study, the inhibition of the MAP-kinase MEK led to the increased anticancer activity of mTOR inhibition both *in vitro* and *in vivo*.<sup>62</sup>

## Reactive Oxygen Species (ROS)

ROS plays a major role in several steps of tumorigenesis.<sup>63</sup> In VHL-defective RCC, ROS production is enhanced because of the impairment of phosphorylative oxidation.<sup>64</sup> It has been demonstrated that PI3K and AKT show increased activity when ROS concentration rises. This response ensures cell survival in stressful conditions.<sup>65</sup>

Another mechanism involves protein kinase C (PKC) family members, which are mTORC2 downstream effectors.<sup>66</sup> The effects of PKC include cell proliferation and epithelial-to-mesenchymal transition by modulation of integrin expression.<sup>67</sup> ROS are able to activate different members of the PKC family. In particular, PKC $\delta$  is activated by ROS in an Src-dependent mechanism that leads to the activation of Jun MAPK.<sup>66</sup>

Finally, it was shown that in RCC, a high concentration of ROS can reduce the degradation of HIF-1 $\alpha$ ,<sup>68</sup> thus activating the aforementioned mechanisms that lead to Rapalogs resistance and cell survival. It appears clear that ROS represent an important mechanism to bypass mTOR inhibition since they can activate directly downstream effectors of the mTOR pathway.

Autophagy/mTOR is considered the principal regulator of the process known as autophagy. Autophagy consists of a mechanism that cells use to recycle damaged organelles in order to recover nutrients and sustain energy production in stressful conditions. It has been demonstrated that mTOR acts as a nutrient sensor, that stimulates cell proliferation, and shuts down autophagy in a ULK1-dependent fashion.<sup>69</sup> Therefore, mTOR blockade acts as a stimulus for autophagy, that causes an increase in nutrients availability to sustain energy production, thus increasing the survival capability of cancer cells.<sup>70</sup> As a counter-proof, available data suggested that the inhibition of the autophagic machinery key component Atg7 can enhance the anti-cancer activity of Rapalogs.<sup>71</sup> From a mechanistic point of view, this process is explained by the impairment in phosphorylative oxidation caused by mTOR inhibition, which leads to an increase in ROS concentration and mitochondrial damage. Damaged mitochondria are usually destroyed via autophagy, but if Atg7 is inhibited, they continue to produce ROS up to the point at which oxidative stress causes cell death. Interestingly, this form of death is not caspase-dependent, being not apoptosis, but rather a form of necroptosis.<sup>71</sup>

## Immune Response Impairment

Rapalogs are primarily considered to be immunosuppressive drugs. Treatment with mTOR inhibitors induces the development of T regulatory cells and makes them more resistant to apoptosis.<sup>72</sup> RCC is an immunologically “hot” tumor and the growing role of immunotherapy in its treatment can be considered proof of the importance of an active immune system in the management of this disease.<sup>73</sup> Data show that in patients treated with Everolimus, the worst prognosis has been demonstrated when the neutrophil to lymphocyte ratio is higher than 3.<sup>74</sup> This is an intrinsic flaw of Rapalogs since their immunosuppressive effect cannot be dissociated from their anticancer effect. However, the relative strength of immune suppression could theoretically be related to the dose used; indeed, the dose commonly administered to transplanted patients for immunosuppressive purposes is completely different from that used in the oncological setting.

On the other hand, mTOR inhibitors could be a suitable option for mRCC in kidney transplant patients, since they could exert both the immunosuppressive and the anticancer effect.<sup>75</sup>

## Overcoming the Resistance to mTOR Inhibitors

The principal clinical trials exploiting strategies to overcome resistance to mTOR inhibitors are summarized in Table 1.

## Second and Third-Generation mTOR Inhibitors

The most immediate strategy to overcome the aforementioned resistance mechanism is the development of more potent mTOR inhibitors. Second-generation mTOR inhibitors, TOR Kinase Inhibitors (TOR-KI), are able to block mTOR in its catalytic domain, by competing with ATP. TOR-KIs do not need FKBP to exert their action since they are developed to bypass this step. The key feature of TOR-KIs, is their ability to block both mTORC1 and mTORC2, thus overcoming some of the aforementioned described mechanisms of resistance. In fact, TOR-KIs were proven to be effective in Rapamycin-insensitive cell lines.<sup>76</sup> However, resistance remains a major concern, since they are not so effective at preventing the feedback activation of PI3K that implies GSK3 and IRS-1.<sup>77</sup>

Vistusertib (AZD2014) was the first molecule of this class to reach clinical experimentation. A phase II clinical

trial compared Vistusertib with Everolimus in patients with mRCC beyond the first line of treatment.<sup>78</sup> The trial was stopped before the completion of the enrollment because of the lack of efficacy observed in Vistusertib's arm. PFS greatly favored Everolimus (1.6 vs 4.8 months,  $p < 0.01$ ). The only true advantage offered by Vistusertib was a slightly more manageable safety profile.

Another TOR-KI that reached human testing is Sapanisertib (TAK-228).<sup>79</sup> A Phase I dose-finding study demonstrated a good safety profile for this drug and an ORR of 22% was observed in one of the cohorts. More clinical trials are needed to point out a possible clinical role for this class of molecules.

Third-generation mTOR inhibitors are currently in their preclinical phase. They were obtained by connecting a first-generation mTOR inhibitor and a second-generation TOR Kinase Inhibitor through a linker.<sup>80</sup> RapaLink-1 is the first compound of this new class, and it is composed of Rapamycin linked to the TOR-KI MLN0128. It was recently compared with Temsirolimus in RCC cell lines, showing a greater anti-tumor efficacy both in Sunitinib-resistant and Sunitinib-sensitive cell lines.<sup>81</sup>

The development of these novel inhibitors could improve the performances observed from the experience with Rapalogs. However, many of the resistance mechanisms described do not depend on the potency of inhibitors per se, but rather on a complex interplay between mTOR and other pathways. Hence the consideration that drugging more than just one molecular target could offer sensible improvements.

## Combination of VEGFR Inhibitors and mTOR Inhibitors

The idea of combining mTOR inhibitors and VEGFR-inhibitors arose soon after the first results of trials with antiangiogenics. Blocking both the angiogenesis and mTOR pathway could deliver enough metabolic damage to RCC cells to improve clinical outcomes and delay the onset of resistance. The first phase I study exploiting this combination was a dose-finding study evaluating the safety of Sunitinib and Everolimus in pretreated mRCC patients.<sup>82</sup> Acute and chronic grade 3 and 4 toxicities were reported among all the cohorts, and the combination resulted to be tolerated only at attenuated doses. The unmanageable safety profile of this combination was confirmed also by a subsequent phase Ib trial that reported similar results.<sup>83</sup>

**Table I** Main Clinical Trials Featuring Strategies to Overcome mTOR Inhibition

Strategy Exploited	Study Population	Study Design	Outcome	Reference
Second-generation mTOR inhibitors	49 mRCC patients that failed a first-line treatment with anti-angiogenic	Phase II trial AZD2014 (2nd generation mTOR inhibitor) vs Everolimus	Everolimus achieved better mPFS (1.8 vs 4.6 months, HR 2.8, p=0.01)	Powles et al. <sup>78</sup>
	82 pretreated patients with mRCC, endometrial or bladder cancer	Phase I trial Dose escalation of Sapanisertib (2nd generation mTOR inhibitor)	ORR 12%, mRCC patients achieved the best responses	Voss et al. <sup>79</sup>
Combination of VEGFR inhibitors and mTOR inhibitors	20 patients with treatment-naïve renal cancer	Phase I trial Dose escalation of Everolimus + Sunitinib	The combination was tolerated, and reported a 25% ORR	Molina et al. <sup>82</sup>
	30 mRCC with 1 ore no previous treatments	Phase I trial Dose escalation of Everolimus + Sunitinib	The combination was not tolerated	Kanesvaran et al. <sup>83</sup>
	153 mRCC patients that failed a first-line treatment with anti-angiogenic	Phase II trial Lenvatinib vs Everolimus vs Lenvatinib + Everolimus	The combination performed better than Everolimus and Lenvatinib in terms of mPFS (14.6 vs 5.5 vs 7.4 months, p1=0.005, p2=0.12)	Motzer et al. <sup>84</sup>
	1069 treatment-naïve mRCC patients	Phase III trial Lenvatinib + Everolimus vs Lenvatinib + Pembrolizumab vs Sunitinib	Lenvatinib + Everolimus performed better than Sunitinib in terms of mPFS (14.7 vs 9.2 months, p<0.001) and it was slightly inferior to Lenvatinib + Pembrolizumab	Motzer et al. <sup>85</sup>
	20 mRCC pretreated patients	Phase I trial Safety evaluation of Vatalanib + Everolimus	The combination was tolerated, and reported a 29% ORR	Bitting et al. <sup>87</sup>
	70 patients with advanced cancer, regardless of the histology	Phase I trial Dose escalation of Vatalanib + Everolimus	The combination was tolerated, and reported a 12.9% ORR and a 58.6% DCR.	Zhu et al. <sup>88</sup>
	22 mRCC pretreated patients	Phase I trial Dose escalation of Vorolanib + Everolimus	The combination was tolerated, and reported a 32% ORR and a 100% DCR.	Sheng et al. <sup>89</sup>
	18 patients with advanced cancer, regardless of the histology	Phase I trial Dose escalation of Vorolanib + Everolimus	The combination was tolerated, and reported a 20% ORR and a 53% DCR.	Pedersen et al. <sup>90</sup>
Alternation of VEGFR inhibitors and mTOR inhibitors	55 treatment-naïve mRCC patients	Phase II trial Alternating Sunitinib and Everolimus	The alternating schedule reported a mPFS of 8 months	Davis et al. <sup>91</sup>
	101 treatment-naïve mRCC patients	Phase II trial Alternating Pazopanib and Everolimus vs continuous Pazopanib	Pazopanib reported a higher mPFS (9.4 vs 7.4 months, p=0.37)	Cirkel et al. <sup>92</sup>
	41 mRCC with 1 ore no previous treatments	Phase II trial Alternating Sunitinib and Everolimus vs standard sequential therapy	Standard sequential therapy reported an higher 1-year PFS (84.6% vs 36%)	Rodriguez-Vida et al. <sup>93</sup>

(Continued)



Table I (Continued).

Strategy Exploited	Study Population	Study Design	Outcome	Reference
PI3K/mTOR dual inhibitors	10 mRCC pretreated patients	Phase I trial Dose escalation of BEZ235 (PI3K/mTOR dual inhibitor)	Study closed due to dose-limiting toxicities.	Carlo et al. <sup>94</sup>
	85 mRCC patients that failed a first-line treatment with anti-angiogenic	Phase II trial Apatolisib (PI3K/mTOR dual inhibitor) vs Everolimus	Everolimus achieved better mPFS (3.4 vs 6.1 months, HR 2.1, $p < 0.01$ ) and ORR (11.6% vs 7.1%)	Powles et al. <sup>95</sup>
Combination of mTOR inhibitors and HDAC inhibitors	13 pretreated patients with mRCC	Phase I trial Dose escalation of Ridaforolimus + Vorinostat (HDAC inhibitor)	The combination is safe. 2 patients achieved long-lasting disease stabilization	Zibelman et al. <sup>97</sup>
	21 pretreated patients with mRCC	Phase I trial Everolimus + Panobinostat (HDAC inhibitor)	The combination is safe. mPFS 4.1 months	Wood et al. <sup>98</sup>
Combination of autophagy inhibitors and mTOR inhibitors	38 patients with mRCC who received 1–3 previous lines of therapy	Phase I/II trial Everolimus + Hydroxychloroquine (autophagy inhibitor)	The combination is safe. ORR 6%, DCR 67%, PFS > 6 months 45%	Haas et al. <sup>100</sup>

**Abbreviations:** mRCC, metastatic renal clear-cell carcinoma; mPFS, median progression-free survival; ORR, objective response rate; DCR, disease control rate.

Another combination explored was Lenvatinib and Everolimus. In a phase II trial,<sup>84</sup> 153 mRCC patients that failed previous antiangiogenic therapy, were randomized to receive either Lenvatinib, Everolimus, or both in combination. The combination arm reported a longer mPFS than both the monotherapy arms (14.6 months vs 5.5 months for Everolimus and 7.4 months of Lenvatinib alone), although the data were statistically significant only in the comparison with the Everolimus arm (HR 0.40, 95% CI 0.24–0.68;  $p = 0.0005$ ). Moreover, the combination arm reported more grade 3 and 4 adverse events, but in the end, the combination therapy showed an acceptable safety profile.

This combination was also evaluated in a phase III trial involving naïve mRCC patients that were randomized to receive Pembrolizumab plus Lenvatinib, Everolimus plus Lenvatinib or Sunitinib.<sup>85</sup> The Pembrolizumab-Lenvatinib arm performed much better than Sunitinib in terms of mPFS (23.9 months vs 9.2 months;  $p < 0.001$ ), but also the Everolimus-Lenvatinib arm reported a good outcome (14.7 months of mPFS). Both the combination arms showed a slightly higher frequency of adverse events, but they had an overall manageable safety profile. The results

reported in these studies led to the design of a clinical trial exploiting Lenvatinib plus Everolimus as a neoadjuvant treatment aimed at making unresectable RCC resectable. This trial is currently ongoing (NCT03324373).

The combination of Lenvatinib and Everolimus was tested also in a phase II trial enrolling a cohort of 31 patients with uncommon RCC subtypes reporting a ORR of 25.8% and a DCR of 61.3%, with a mPFS of 9.2 months.<sup>86</sup> This is an important milestone for the treatment of RCC variants that often lack of valuable therapeutic options.

Some better results come from a phase Ib trial of the VEGFR-inhibitor Vatalanib in combination with Everolimus.<sup>87</sup> Even though adverse events were observed, long-term tolerability was demonstrated in a significant portion of patients. Moreover, an mOS of 16.3 months was reported with an ORR of 29.2%. Interestingly, the mOS was much longer for treatment-naïve patients (25.1 months). These data suggest that the combination of Vatalanib and Everolimus could be more effective than the sequential schedule at obtaining long-lasting responses. More recently, another phase I trial analyzed this combination in a cohort of metastatic solid tumors, and the data were confirmed in the mRCC subgroup.<sup>88</sup>

Similar results were reported by a phase I trial investigating the combination of the VEGFR-inhibitor Vorolanib (X-82, CM082) and Everolimus in mcrRCC pretreated patients.<sup>89</sup> Along with the good tolerability of the treatment, a good efficacy emerged from this study, with an ORR of 32% and a DCR of 100%. These data are confirmed also by another phase I trial exploiting this combination in solid tumors: the mcrRCC cohort was the one with the best ORR.<sup>90</sup>

In the end, the combination of VEGFR inhibitors and mTOR inhibitors resulted to be difficult to manage in some cases but reported an efficacy profile good enough to prompt further investigation.

## Alternating VEGFR Inhibitors and mTOR Inhibitors

A theoretically simple strategy to avoid the toxicities of the combination, while reducing the resistance phenomena, could be the sequential alternation of VEGFR inhibition and mTOR inhibition. The rationale of this therapeutic schedule resides in the hypothesis that VEGFR-inhibitors would select a clone of RCC cells that are sensitive to mTOR inhibition, and vice-versa. The EVERSUN phase II trial tested an alternating schedule of Sunitinib and Everolimus in 55 naïve mcrRCC patients, obtaining a median PFS of 8 months and a median OS of 17 months.<sup>91</sup> These results did not meet the pre-specified values to warrant further research.

In the ROPETAR trial, 101 naïve mcrRCC patients were randomized to receive either an alternating schedule of Pazopanib and Everolimus or continuous Pazopanib.<sup>92</sup> The Pazopanib arm performed better in terms of median PFS (9.4 vs 7.4 months) and no significant differences were observed in the safety profile, thus suggesting that the rotating schedule gives no advantages at all.

Similar findings were reported by the SUNRISE study, a phase II trial comparing alternating cycles of Sunitinib and Everolimus versus standard sequential treatment of Sunitinib followed by Everolimus upon progression, in 42 naïve mcrRCC patients.<sup>93</sup> The 1-year PFS rate was 49.7% in the alternating arm and 84.62% in the sequential arm, with no major differences in mOS. From this data, it can be inferred that the alternating strategy fails to improve the clinical outcomes in mcrRCC. The almost contemporary exposure to both drugs likely favors the development of resistance mechanisms allowing mcrRCC cells to acquire resistance to both VEGFR-inhibition and mTOR-inhibition.

## PI3K/mTOR Dual Inhibitors

Among all the resistance mechanisms analyzed, the PI3K feedback is surely one of the most common, and its onset cannot be delayed by simply increasing the potency of mTOR blockade. Hence the idea of developing molecules that can block both mTOR and PI3K.

One of the first clinical studies involving this new class of drugs was a phase Ib trial of BEZ235 previously treated patients with mcrRCC.<sup>94</sup> Even though BEZ235 was extensively tested in the preclinical setting, the trial was interrupted due to the high incidence of dose-limiting toxicities including fatigue, rash, nausea and vomiting, diarrhea, mucositis, anorexia, and dysgeusia. Therefore, the development of the drug was stopped.

Another drug of this class, Apatolisib (GDC-0980) was compared with Everolimus in a phase II clinical trial enrolling mcrRCC patients that progressed on after the first-line treatment with VEGFR-inhibitors.<sup>95</sup> Surprisingly, the Everolimus arm had a longer median PFS (6.1 vs 3.7 months) and median OS (22.8 vs 16.5 months), and a higher ORR (11.6% vs 7.1%). The authors explained these results with the higher frequency of adverse events observed in the Apatolisib arm. A possible explication for the unexpected bad performance of PI3K/mTOR dual inhibitors could be that the resistance mechanisms implying PI3K, although frequent, are not pivotal for cell survival. Moreover, the unfavorable safety profile of these drugs makes it difficult to evaluate objectively their results. Finally, it should be pointed out that the design of this trial could have been improved. In fact, Apatolisib could have been administered to patients progressed beyond a first line therapy with anti-angiogenics and a second line therapy with Everolimus. In this setting, Apatolisib should have been compared with best supportive care or clinician choice, thus demonstrating if it is effectively capable of overcoming the resistance built by Everolimus through its innovative mechanism of action.

## Combination of mTOR Inhibitors and Histone Deacetylase (HDAC) Inhibitors

The inhibition of HDAC exerts its anticancer effect by modulating DNA coiling and thus, genes expression. Preclinical data showed that AKT phosphorylation is negatively affected by HDAC inhibition, although the precise mechanism of the phenomenon remains unclear.<sup>96</sup> Hence the rationale for combination therapy.

A recent phase I clinical trial tested the rapalog Ridaforolimus with the HDAC inhibitor Vorinostat in mRCC pretreated patients.<sup>97</sup> Although it was only a dose-finding study, some of the patients experimented a long-lasting disease stabilization, and no safety concerns were observed. Similar results were observed in another phase I trial with the combination of Everolimus and the HDAC inhibitor Panobinostat.<sup>98</sup> Moreover, this combination reported a 4.1 months median PFS, although more standardized studies involving larger populations are needed to clarify the possible role of this combination in mRCC therapy.

## Combination of Autophagy Inhibitors and mTOR Inhibitors

As discussed before, mTOR blockade promotes autophagy processes that cancer cells use to recycle damaged organelles and produce energy. The drug Hydroxychloroquine is known for its ability to inhibit autophagy, by limiting the fusion of lysosomes and autophagosomes.<sup>99</sup> Haas et al studied the efficacy and safety of combining Hydroxychloroquine and Everolimus in a cohort of 33 patients that had already failed previous therapy lines. Disease control was achieved in 66% of patients, with an ORR of 6%. No dose-limiting toxicity was reported. Interestingly, this phase I/II trial met its primary endpoint, with a 6-months-PFS higher than 40%.<sup>100</sup> The data shown here confirm the importance of autophagy in the onset of resistance to rapalogs, but further investigation is necessary to understand exactly the extent of the improvement offered by autophagy inhibitors.

## Conclusion

Despite the rationale of targeting mTOR in mRCC, and its pivotal role in RCC tumorigenesis, the inhibition of mTOR through Rapalogs did not meet the expectations. Objective responses were scarce and, more importantly, disease control was brief.

These results are principally attributable to the complex interplay between the mTOR pathway and many other cell growth and proliferation pathways, that allows the onset of resistance phenomena. At the state of the art, some solutions to overcome the resistance are under development. On the one hand, more potent and innovative mTOR inhibitors are being created, to obtain an improved blockade of both mTORC1 and mTORC2. On the other hand, combinatorial strategies aim at

countering some of the known resistance mechanisms. In the near future, more combinations of mTOR inhibitors and VEGFR inhibitors could reach clinical practice. More studies are needed for combinations including autophagy blockers and HDAC inhibitors, both of which showed promising preliminary results.<sup>97,98,100</sup> Finally, completely unexplored remains the combination of mTOR inhibitors and immune checkpoint inhibitors.

Even if it may appear counterintuitive (potentially immune suppressive agents combined with an immune stimulating one?), this synergy could improve the effectiveness of immune therapy, which has already shown excellent results in mRCC.

## Disclosure

Dr Mimma Rizzo reports personal fees from MSD, personal fees from Pfizer, personal fees from Novartis, personal fees from AstraZeneca, during the conduct of the study.

Prof. Dr. Camillo Porta reports personal fees from Angelini Pharma, personal fees from AstraZeneca, personal fees from BMS, personal fees from Eisai, personal fees from EUSA Pharma, personal fees from General Electric, personal fees from Ipsen, personal fees from Janssen, personal fees from Merck Serono, personal fees from MSD, personal fees from Novartis, personal fees from Pfizer, and Roche, outside the submitted work. The authors report no conflicts of interest relative to the present manuscript.

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