The c-Met Tyrosine Kinase Inhibitor JNJ-38877605 Causes Renal Toxicity through Species-Specific Insoluble Metabolite Formation

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Abstract

Purpose: The receptor tyrosine kinase c-Met plays an important role in tumorigenesis and is a novel target for anticancer treatment. This phase I, first-in-human trial, explored safety, pharmacokinetics, pharmacodynamics, and initial antitumor activity of JNJ-38877605, a potent and selective c-Met inhibitor.

Experimental Design: We performed a phase I dose-escalation study according to the standard 3 + 3 design.

Results: Even at subtherapeutic doses, mild though recurrent renal toxicity was observed in virtually all patients. Renal toxicity had not been observed in preclinical studies in rats and dogs. Additional preclinical studies pointed toward the rabbit as a suitable toxicity model, as the formation of the M10 metabolite of JNJ-38877605 specifically occurred in rabbits and humans. Additional toxicology studies in rabbits clearly demonstrated that JNJ-38877605 induced species-specific renal toxicity.

Introduction

Tumor cell migration, invasiveness, and metastasis formation are important therapeutic targets in oncology drug development. The hepatocyte growth factor (HGF), also known as scatter factor, and its receptor c-Met, play an important role in physiologic processes such as embryologic development, wound healing, tissue regeneration, angiogenesis, cell growth, local invasion, and morphogenetic differentiation (1). In physiologic and pathophysiologic situations, HGF is primarily produced by fibroblast-like cells, whereas c-Met is expressed by epithelial cells. Aberrant expression of c-Met is involved in tumor progression (2, 3). Activating c-Met mutations have been identified in hereditary and sporadic papillary kidney cancer and gastric carcinomas, suggesting a potential causal role for c-Met activation in tumorigenesis (4, 5). Moreover c-Met amplification is a well-known event in non–small cell lung cancer especially important in resistance to EGFR inhibition (6). In a small subset of glioblastoma multiforme patients, activating c-Met mutations have been reported and activation of the HGF–c-Met axis seems to play a role in resistance to therapy (7). Therefore, inhibiting the activity of c-Met is a relevant objective in anticancer drug development.

JNJ-38877605 is an orally available, nanomolar active (IC50 4.7 nmol/L for c-Met kinase in vitro) and highly selective c-Met ATP-competitive kinase inhibitor (8). JNJ-38877605 inhibits c-Met kinase with >333-fold selectivity relative to the next most potently inhibited kinase (Fms) of 246 kinases tested. JNJ-38877605 binds to the ATP-binding site of c-Met kinase with a high affinity that leads to a slow reversibility of binding. JNJ-38877605 has shown a favorable safety profile in rat and dog studies and demonstrated antitumor activity in xenograft models of prostate, non–small cell lung and gastric cancer as well as in glioblastoma models.

This phase I, open-label, dose-escalation study explored safety, tolerability, and pharmacokinetics of orally administered JNJ-38877605.

Materials and Methods

Patients and eligibility criteria

This was a nonrandomized, open-label, phase I, dose-escalation study conducted at two participating institutions...
Translational Relevance

The early clinical testing of novel target-specific agents is an important part of improving anticancer treatment. Preclinical animal data guide initial clinical safety assumptions, and dogs, mice, and rats are the most frequently used species. This first-in-human study with JNJ-38877605 revealed renal toxicity, even at subtherapeutic doses, that was not predicted by the aforementioned animal models. Subsequently performed rabbit studies showed that species-specific metabolism of quinoline ring structures of JNJ-38877605 through aldehyde oxidase resulted in the formation of insoluble metabolites that form crystal structures in the kidney leading to renal toxicity. Our findings suggest that in case of involvement of aldehyde oxidase, rabbit models should be used as alternative toxicology species.

Pharmacokinetics

Blood samples were taken at prespecified times (see Supplementary Methods) and analyzed quantitatively for [N]-38877605 and the N-desmethyl metabolite, M2. For both compounds, Cmax time to Cmax (tmax), elimination half life time (t1/2), area under the plasma concentration curve (AUC), and terminal elimination rate constant were calculated.

Toxicology studies in the model organisms

Toxicology studies were performed according to ICH Topic S9. Nonclinical Evaluation for Anticancer Pharmaceuticals, EMEA/CHMP/ICH/646107/2008, and ICH guideline M3(R2) on nonclinical safety studies for the conduct of human clinical trials and marketing authorization for pharmaceuticals. Initially male and female rats and dogs were chosen as toxicology species. The choice of the toxicology species is based on a comparison of the metabolism pattern in hepatocytes of preclinical species and man. Human metabolites should be covered in the toxicology species to assess compound and metabolite-related toxicity during the 1-month GLP toxicology study. On the basis of metabolite patterns in hepatocytes, all human metabolites were present in rat and dog with the exception of an N-glucuronide, which is considered to be nontoxic. Therefore, rat and dog were identified as suitable species to test toxicity.

Following the observation of renal toxicity in the clinical study, we chose rabbits because of the presence of the glucuronide of the desmethyelmetabolite (M10) in rabbit and man, only. A 1-month non-GLP toxicology study of JNJ-38877605 was performed in the male rabbit and histopathological evaluation of the kidney was performed as well as all other standard tissue testing and sampling. To find possibilities to circumvent renal toxicity, various intermittent dosing schedules were explored (see Table 1 for the dosing regimen and total weekly dose).

Metabolite identification

Metabolites were determined from in vitro systems (hepatocytes and liver subcellular fractions), and in vivo samples (plasma, urine, and kidney samples). Metabolite identification was performed by LC/MS, quantitative data are only available for those metabolites with a reference standard or metabolites with a 3H label. Solubility measurements were performed for some of the metabolites.

Isoenzymes involved in the formation of the metabolites

Isoenzyme identification was performed with liver subcellular fractions. Isoenzymes involved in the metabolism of JNJ-38877605 were determined based on structural characteristics and after incubation with specific inhibitors of cytochrome P-450 and aldehyde oxidase (9).

Histopathological evaluation

Rabbit kidneys were fixed by formalin immersion, then processed routinely: trimming, embedding, sectioning at 3 to 5 μm, staining with hematoxylin and eosin (H&E), and examined under bright field. Additional frozen sections of kidney were prepared and counterstained with either methylene blue or Von Kossa (to detect mineralization). They were examined under bright field and polarization microscopy.
Identification of the crystal structures in the kidneys
Kidney crystals detected under polarization microscopy were observed and analyzed in situ using Raman microscopy. Therefore, they were harvested and analyzed with LC/MS.

Prevention of renal toxicity
A mechanistic study was performed to investigate the potential protective effect of probenecid on renal toxicity. Probenecid is known as a competitive inhibitor of organic acid transport in the kidney and other organs (10). After exposure to JNJ-38877605 in combination with probenecid, determination of pharmacokinetics (plasma, kidney, urine) of JNJ-38877605, renal parameters (blood urea nitrogen and creatinine), and kidney histopathology were performed in male rabbits.

Results
Toxicity in the phase I, first-in-human study of JNJ-38877605
A total of five subjects were enrolled in the first cohort of 60 mg OD. All subjects experienced a rapid, though asymptomatic grade 1 serum creatinine (SCr) increase; in 4 patients this occurred as early as 2 to 3 days after the start of treatment, whereas 1 patient completed 4 weeks of treatment before SCr increase. In all cases, treatment was interrupted. In 1 patient, a rechallenge with JNJ-38877605 at the same dose was pursued after a 3-day interruption of therapy and subsequent normalization of SCr; this patient again experienced a grade 1 SCr increase after which treatment was permanently discontinued. All subjects recovered from their SCr increase after permanent treatment discontinuation.

To explore potential safe dose levels, an amendment was pursued after a 3-day interruption of therapy and subsequent normalization of SCr; this patient again experienced a grade 1 SCr increase after which treatment was permanently interrupted. Due to the rapid and transient nature of these events, this patient received no further treatment. All subjects recovered from their SCr increase after permanent treatment discontinuation. In 1 patient, a rechallenge with JNJ-38877605 at the same dose was pursued after a 3-day interruption of therapy and subsequent normalization of SCr; this patient again experienced a grade 1 SCr increase after which treatment was permanently interrupted. All subjects recovered from their SCr increase after permanent treatment discontinuation.

Pharmacokinetic analysis of JNJ38877605 in humans
JNJ38877605 was rapidly absorbed and showed a Cmax at 1.5 to 4 hours after ingestion (Fig. 1C). A short plasma half-life and virtually no accumulation after 21 days were found. Comparison of plasma concentrations and AUC values over the different dosing cohorts showed dose proportionality (Supplementary Table S2). At 60 mg OD, plasma concentrations and AUC did not reach levels that in preclinical models resulted in antitumor activity.

Interspecies difference in metabolism of JNJ-38877605
On the basis of a previous interspecies comparison of its metabolism, it was hypothesized that the renal toxicity of JNJ-38877605 that already occurred at subtherapeutic doses, could possibly be explained by species-specific metabolism. After incubation of JNJ-38877605 with liver fractions and identification of the metabolites with LC/MS-MS, several metabolites were observed. The major metabolic pathways in humans were demethylation, leading to M2, which was further metabolized to M5/M6 by hydroxylation of the quinoline moiety and to M10 by glucuronidation. JNJ-38877605 was also hydroxylated at the quinoline moiety to M1/M3. (Fig. 2 and Supplementary Table S3). A comparative metabolism study in hepatocytes of rat, dog, rabbit, and man showed a species-dependent formation of...
M10 (major in man and rabbit) and M11 (minor; Supplementary Fig. S1). We therefore hypothesized that M10 could play a role in the observed clinical renal toxicity and subsequently selected rabbit as a toxicity model.

Renal effects after JNJ-38877605 administration during 1 month in rabbits

After total weekly intake of 200, 300 and 350 mg/kg/week for 1 month, creatinine clearance and several other chemistry parameters started changing. At necropsy (Table 1), rabbit kidneys showed increased weights (groups 3–6) and appeared pale (groups 4–6). Histopathological evaluation showed an extensive and prominent degeneration with inflammation, congestion, fibrosis, regeneration, and granular and acicular clefts sometimes surrounded by giant cells (Table 1; Fig. 3B–D). In frozen sections examined under polarization microscopy, these clefts proved to be bi-refringent crystalline granules and spicules (Fig. 3E and F). Some of the observed changes, in particular the kidney weight changes and the crystal accumulation, were still present after a 1-month recovery period (Supplementary Table S4), with a moderate increase in urea nitrogen and a slight increase in creatinine.

As renal crystal formation was probably related to the formation of insoluble metabolites of JNJ-38877605 in the urine, we explored the potential protective effect of coadministration of probenecid. As probenecid is a competitive inhibitor of organic acid transport in the kidney and other organs, we hypothesized that it could block the renal transport of JNJ-38877605 and its metabolites. In a 7-day rabbit study, probenecid coadministration, however, resulted in more severe clinical observations compared with dosing with JNJ-38877605 alone and in similar histopathological changes (Supplementary Table S5). In addition, exposure to JNJ-38877605 decreased after probenecid coadministration (data not shown).

Identification of the JNJ-38877605 derived metabolites in the renal crystals

We next harvested crystals from open slides without coverslips and analyzed those with LC/MS; two chemical structures were proposed based on LC/MS results. Additional comparison of the LC-mass UV data with TOX LC-mass UV data and UV reference spectra allowed for the conclusion that both proposed structures contain a 2-hydroxyquinoline chromophore, with proposed
structure 1 relating to JNJ-38877605 metabolites M5, M6 and proposed structure 2 relating to metabolites M1, M3. Additional experiments showed that the M5 metabolite displayed poor solubility at with a maximum concentration of 2 to 3 mg/mL solute over a range of pH.

We next performed mass balance studies in rats, dogs, and rabbits showing that the excretion of JNJ-38877605 and its metabolites occurs partly through the kidney and rabbits and humans predominantly excrete M3 and M5 in the urine. In rabbit, urinary excretion was more important compared with rat and dog (data not shown). Plasma profiling showed a much lower abundance of M3 and M5 in rat and dog compared with rabbit and human (Table 2). These data suggested that plasma levels of M3 and M5 and their renal excretion play a critical role in the explanation of the renal findings.

Identification of the isoenzymes involved

Identification of the isoenzymes involved in the metabolism of JNJ38877605 is based on studies with specific inhibitors in liver microsomes and supernatant fractions. M3 is the most important metabolite formed after incubation of JNJ-38877605 with human liver supernatant and is almost undetectable in human liver microsomes (Table 3). This suggests that M3 is primarily formed by cytosolic enzymes. This is further corroborated by the fact that quercetin completely inhibited the formation of M3 in 12,000 × g fractions (Table 3). Quercetin is a potent inhibitor of molybdenum hydroxylases like xanthine oxidase and aldehyde oxidase. Aldehyde oxidase is known to oxidize quinoline moieties, therefore most probably aldehyde oxidase and not xanthine oxidase is involved in the metabolism of JNJ-38877605. In addition, 1-aminobenzotriazole, a general cytochrome P-450 inhibitor, barely inhibits the M3 formation which is in line with the assumptions above (Table 3). Cytochrome P-450 3A4 is clearly involved in the formation of the other metabolites. M5, the equivalent of M3 but originated from the demethylated JNJ-38877605 (M2) was not formed in microsomes or 12,000 × g. On the basis of the structural similarity with M3, it is assumed that it is also formed by aldehyde oxidase.

Discussion

In this first-in-human phase I study with the c-Met tyrosine kinase inhibitor JNJ-38877605 renal toxicity, hampering...
continuation of treatment, was observed after a short timeframe and at virtually all, even subtherapeutic dose levels. Based upon the at that time available results from preclinical studies performed in mice, rats, and dogs, this renal toxicity had not been foreseen. Extensive additional preclinical research therefore was initiated. On the basis of the subsequent analysis of metabolism data, the rabbit was subsequently identified as a relevant toxicity model. In this model, we were able to recognize aldehyde oxidase-dependent and species-specific formation of insoluble metabolites that induced renal toxicity through precipitation of insoluble crystals in the renal tubules.

In the clinical study, although strongly and repeatedly considered, it was decided not to perform renal biopsies in any of the patients, and therefore the presence of such crystals in any of our patients could not be confirmed. This decision was based upon the clinical and ethical consideration that patients in this study had only asymptomatic grade 1 SCr increases.

Unfortunately, we were not able to come up with a viable strategy to circumvent renal toxicity in the subsequently performed rabbit studies, and therefore it was decided that further clinical development of JNJ-38877605 had to be stopped.

Table 2. Metabolites of JNJ-38877605 in plasma

<table>
<thead>
<tr>
<th>Species</th>
<th>M1</th>
<th>M2</th>
<th>M3</th>
<th>M4</th>
<th>M5</th>
<th>M6</th>
<th>M7</th>
<th>M10</th>
<th>UD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rat</td>
<td>136</td>
<td>64</td>
<td>57</td>
<td>15</td>
<td>6</td>
<td>1,466</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dog</td>
<td>31</td>
<td>55</td>
<td>40</td>
<td></td>
<td></td>
<td>575</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rabbit</td>
<td>6,388</td>
<td>778</td>
<td>603</td>
<td>2,259</td>
<td>1,780</td>
<td>395</td>
<td>822</td>
<td>22,836</td>
<td></td>
</tr>
<tr>
<td>Man</td>
<td>387</td>
<td></td>
<td>208</td>
<td></td>
<td></td>
<td>2</td>
<td>111</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

NOTE: The metabolites of JNJ-38877605 were determined in urine/feces of rat, dog, and rabbit and in urine of man. Animals were dosed with 3H-JNJ-38877605 at 2.5 mg/kg in rat, 2.5 mg/kg in dog, and 100 mg/kg in rabbit. Humans were dosed at 60 mg. JNJ-38877605 and its metabolites in plasma were expressed as ngeq/mL. Metabolites were measured at about Cmax, ranging between 1 hour and 4 hours depending on the species.

Abbreviation: UD, unchanged drug or JNJ-38877605.
With regard to the observed clinical toxicity of JNJ-38877605, it is intriguing that seemingly comparable observations were made in a phase I study with SGX523, another ATP-competitive small-molecule c-Met inhibitor (11). At doses exceeding 80 mg, renal toxicity consisting of early rises in BUN and creatinine and obstructive nephropathy were observed. SGX523 equally showed toxicity consisting of early rises in BUN and creatinine and which also prompted investigators to interrupt further clinical development (11, 12). These data underpin our observation that quinoline-containing chemical structures can be metabolized in a species-specific manner to form nephrotoxic poorly soluble metabolites explaining the observed renal toxicity of SGX523 and JNJ-38877605.

In conclusion, JNJ-38877605 is an orally available, nanomolar inhibitor of the metabolism of JNJ-38877605 and the formation of the major metabolites in human liver supernatant by an inhibitor of aldehyde oxidase and cytochrome P-450. 

**Disclosure of Potential Conflicts of Interest**

H.H. Bohets, A. Lampo, E. Barale, and P. Hellemans have ownership interest in Johnson & Johnson. H.H. Bohets, A. Lampo, E. Barale, M.J.A. Dejonge, L. van Doorn, P. Hellemans, J. deBono, F.A.L.M. Eskens, M.P. Lolkema, E. Barale, M.J.A. Dejonge, L. van Doorn, have ownership interest in Johnson & Johnson. J. DeBono is a consultant/advisory board member for Johnson & Johnson. No potential conflicts of interest were disclosed by the other authors.

**Authors’ Contributions**

Conception and design: H.H. Bohets, A. Lampo, P. Hellemans, J. deBono, F.A.L.M. Eskens

Development of methodology: H.H. Bohets, A. Lampo, E. Barale, J. deBono, F.A.L.M. Eskens

Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): M.P. Lolkema, E. Barale, M.J.A. Dejonge, L. van Doorn, P. Hellemans, J. deBono, F.A.L.M. Eskens

**Clinical and Preclinical Assessment of TKI JNJ-38877605**

**Table 3. Identification of the enzymes causing generation of the insoluble metabolite M3**

<table>
<thead>
<tr>
<th>Matrix</th>
<th>Metabolites (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Microsomes</td>
<td>M1/M2 M3 M4 UD</td>
</tr>
<tr>
<td>12,000 &lt; g</td>
<td>4.46 1.21 4.07 12.8</td>
</tr>
</tbody>
</table>

5 μmol/L TH-JNJ-38877605 was incubated with human liver microsomes or 12,000 < g during 120 minutes.

**Inhibition of the metabolism of JNJ-38877605 and the formation of the major metabolites in human liver supernatant by an inhibitor of aldehyde oxidase and cytochrome P-450**

<table>
<thead>
<tr>
<th>Inhibitor</th>
<th>Enzyme</th>
<th>% inhibition of metabolite formation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Quercetin</td>
<td>Aldehyde oxidase</td>
<td>75 94 73</td>
</tr>
<tr>
<td>1-aminobenzotriazole</td>
<td>CYP-450</td>
<td>58 1 70</td>
</tr>
</tbody>
</table>

NOTE: 5 μmol/L TH-JNJ-38877605 was incubated with human liver supernatant. Abbreviation: UD, unchanged drug or JNJ-38877605.
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): M.P. Lolkema, H.H. Bobets, J. deBono, F.A.L.M. Eskens

Writing, review, and/or revision of the manuscript: M.P. Lolkema, H.H. Bobets, E. Barale, M.J.A. Dejonge, J. deBono, F.A.L.M. Eskens

Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): H.H. Bobets, E. Barale, P. Hellemans, J. deBono, F.A.L.M. Eskens

Study supervision: M.P. Lolkema, H.H. Bobets, E. Barale, P. Hellemans, J. deBono, F.A.L.M. Eskens

Acknowledgments

The authors thank all the patients, nurses, and site staff who participated in the study, the compound development leader Peter Palmer, all collaborators of the preclinical toxicology, PDMS and pharmacokinetics departments for their dedication to the project, and more specifically Tom Peeters, Petra Vinken, Sofie Starckx, Dirk Cleerens, John Vandenbergh, Loes Versmissen, Kristel Buyens, Hilde De Man, Katelijne Anciaux, Ellen Scheers, Carine Borgmans, Jan Snoey, Sandy Thijsen, Rudy van Beijsterveldt, and Brendan Rooney.

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