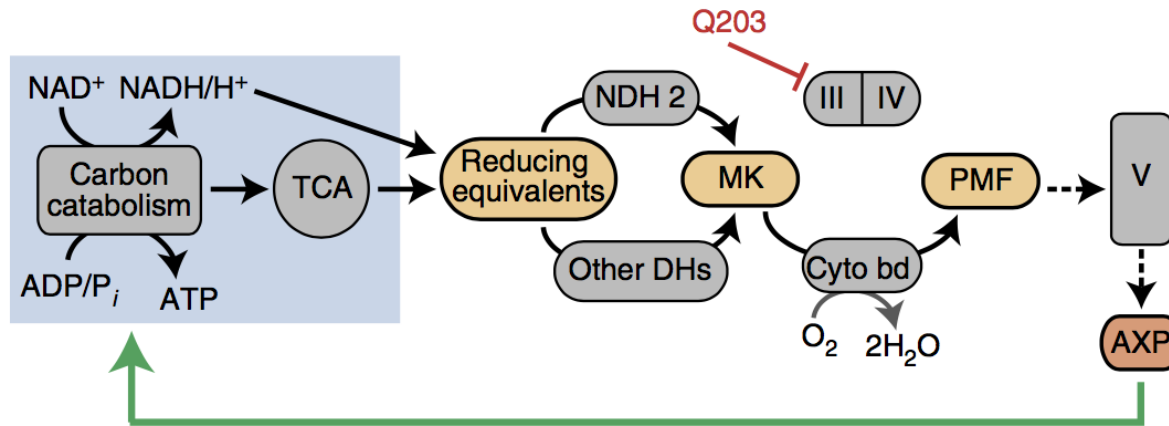


Qurient

Therapeutics

Q203 Update

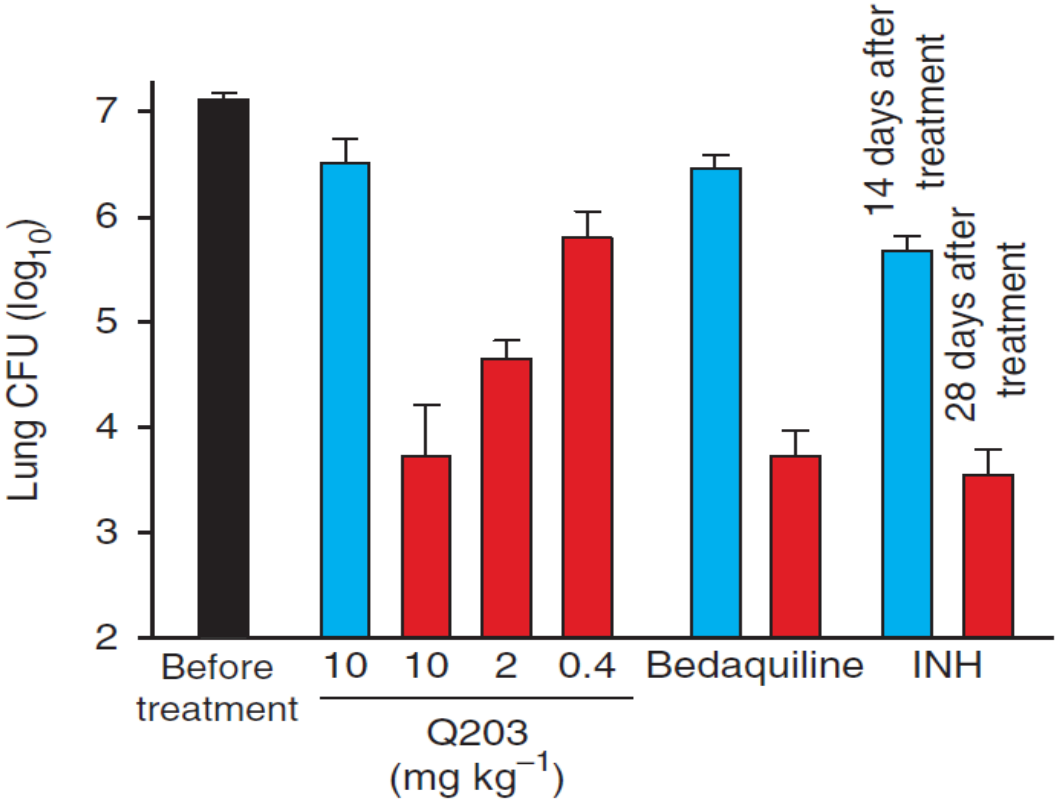
- Bacteria killing by
Inhibition of respiratory chain (cytochrome bc1 complex)



- Host anti-inflammation by
Inhibition of 5 lipoxygenase

Q203: Bacteria Killing Activity

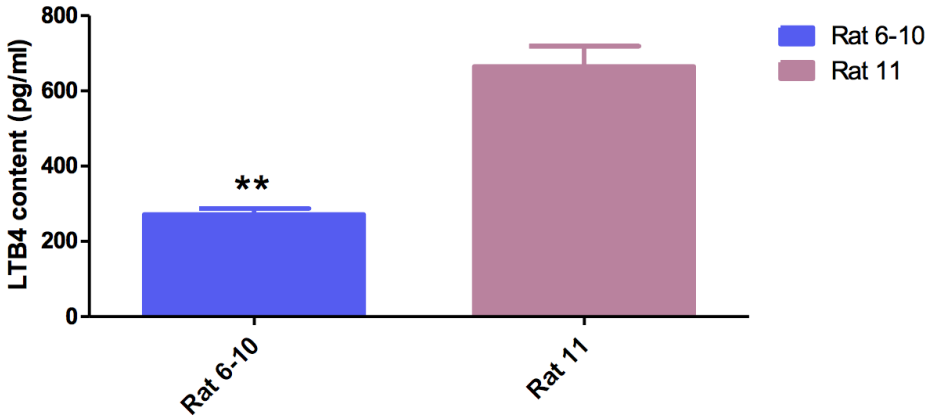
- Mouse 4 weeks treatment model shows good efficacy
- Maximum efficacious dose at 10 mg/kg



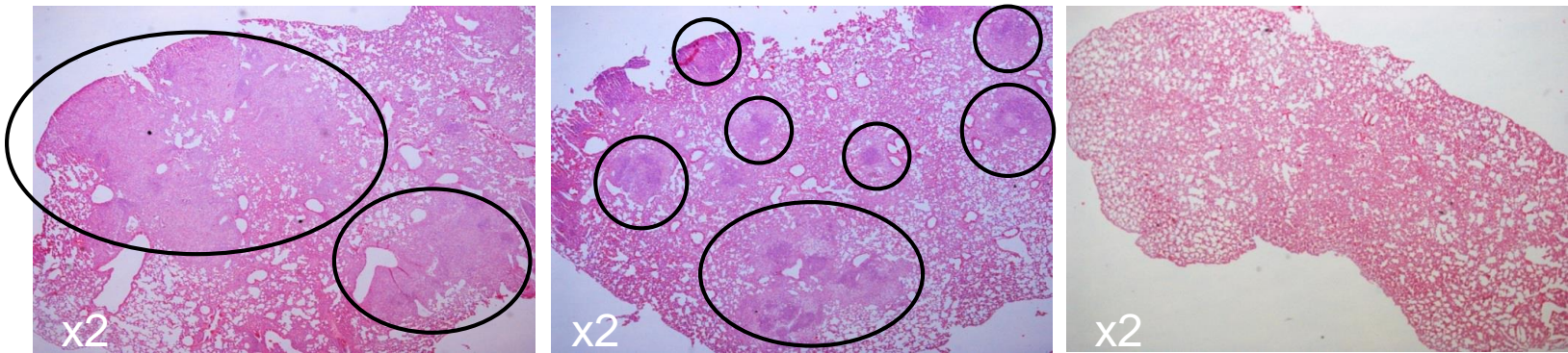
(Nature Medicine 2013)

Q203: Anti-Inflammation

- Oral administration of Q203 inhibits production of LTB4



- Reduction in numbers of granuloma in mouse TB model



Untreated (vehicle)

INH (15 mg/kg)

Q203 (10 mg/kg)

(Nature Medicine 2013)

- Absorption

Mouse single dose in fasted condition shows long half life (~90 hour)

Steady-state after 14 days of dosing

- Distribution

Mouse mass balance study shows compound distribution to lung

- Metabolism

Negative DDI sign from in vitro study

No major metabolite

- Excretion

Mainly excreted through feces

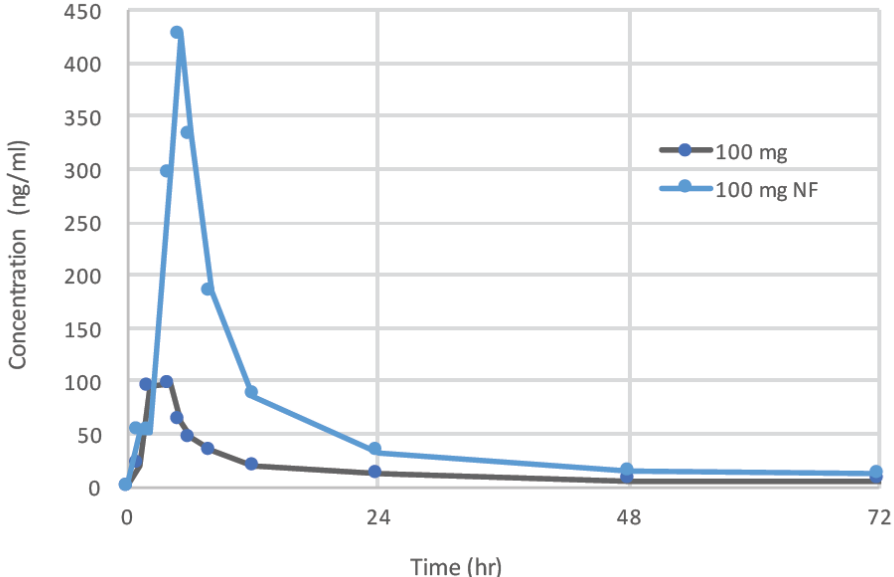
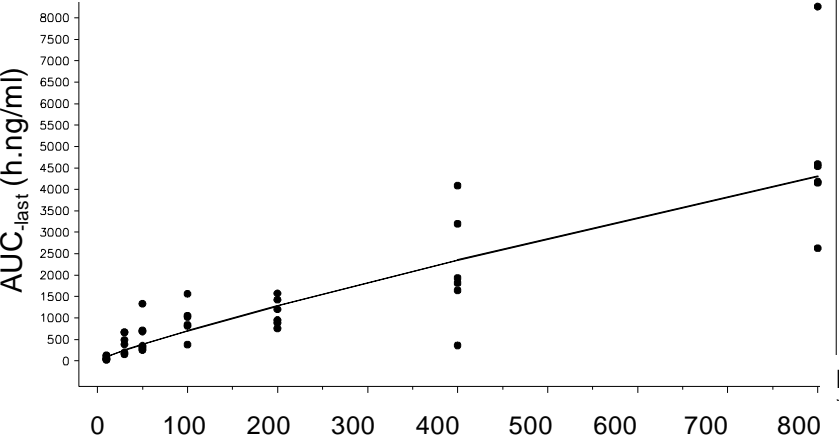
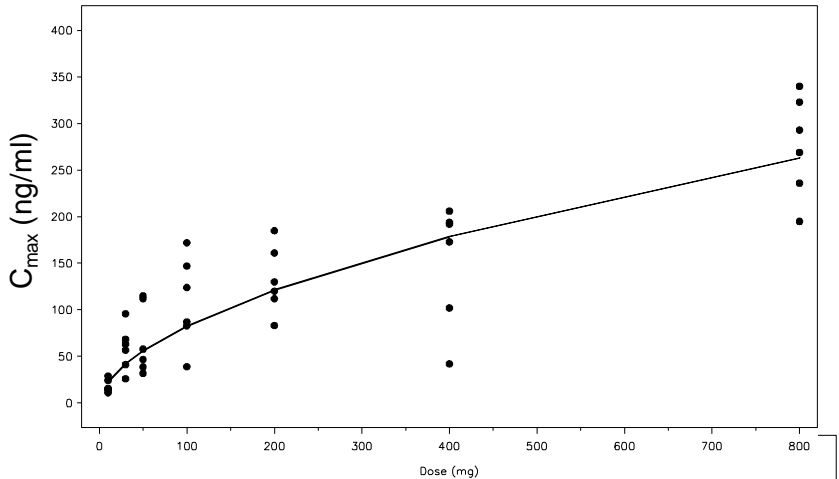
- Toxicity

Well established NOAEL up to 28 days

- Absorption
 - Mouse single dose in fasted condition shows long half life (~90 hour)
 - Steady-state after 14 days of dosing
- Distribution
 - Mouse mass balance study shows compound distribution to lung
- Metabolism
 - Negative DDI sign from in vitro study
 - No major metabolite
- Excretion
 - Mainly excreted through feces
- Toxicity
 - Well established NOAEL up to 28 days

Q203: Phase 1 SAD

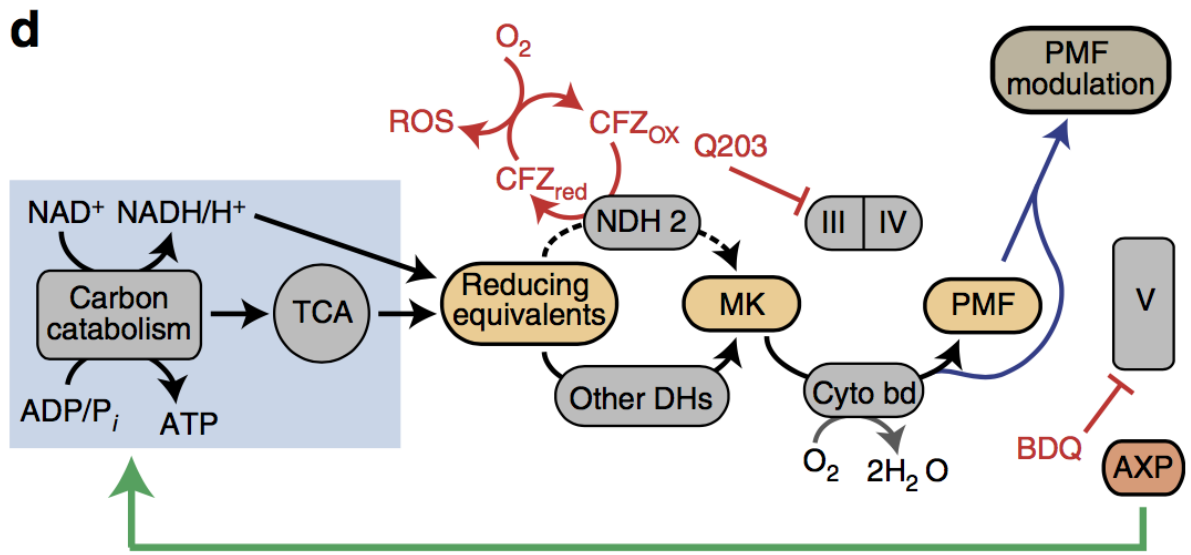
- Dose proportional increase of C_{max} and AUC_{last}
- No clinically significant changes in safety parameters including ECG
- Food increases absorption around 5 fold



- Designed to support 14-day EBA study
- Dose escalation up to maximum anticipated efficacious level

Q203: Future Plan

- EBA study planned starting end of 2017
Standard 14-day EBA with extended combination dosing phase with HRZ or innovative backbone (bedaquiline ?)



- Developing safety and efficacy profile in combination regimen from early stage

