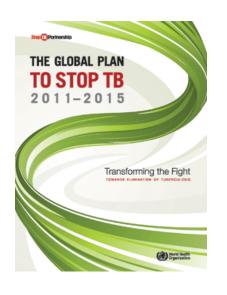


Progress Report 2011

Mel Spigelman, Co-Chair

Mission

To accelerate the discovery and development of new tuberculosis treatments by bringing together all stakeholder perspectives, including those of the patients, in TB Drug Research and Development



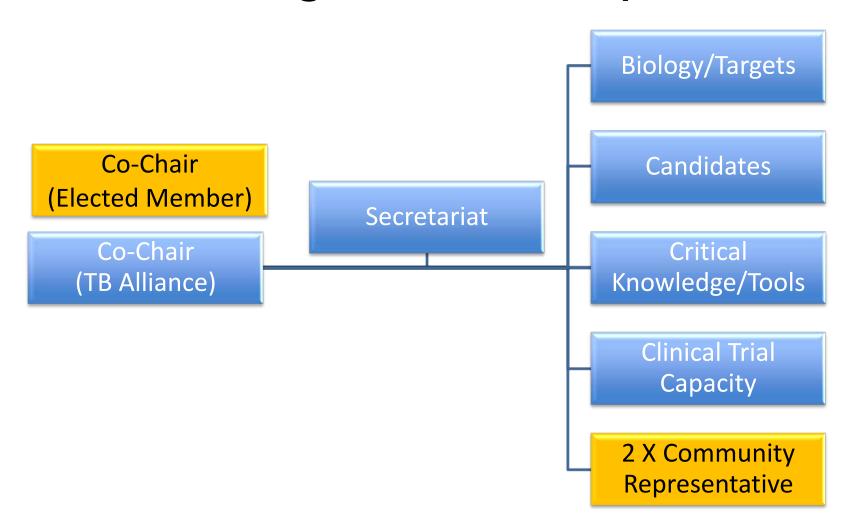


- Resource Gateway
- Catalyst of Dialog and Exchange
- Dynamic and Expanding Network



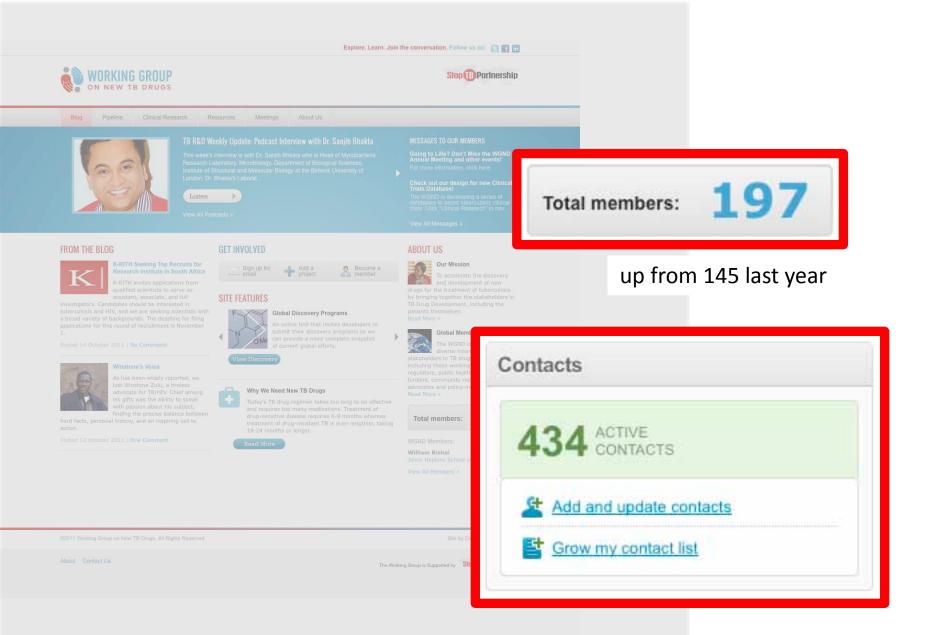


WGND Organization Updates









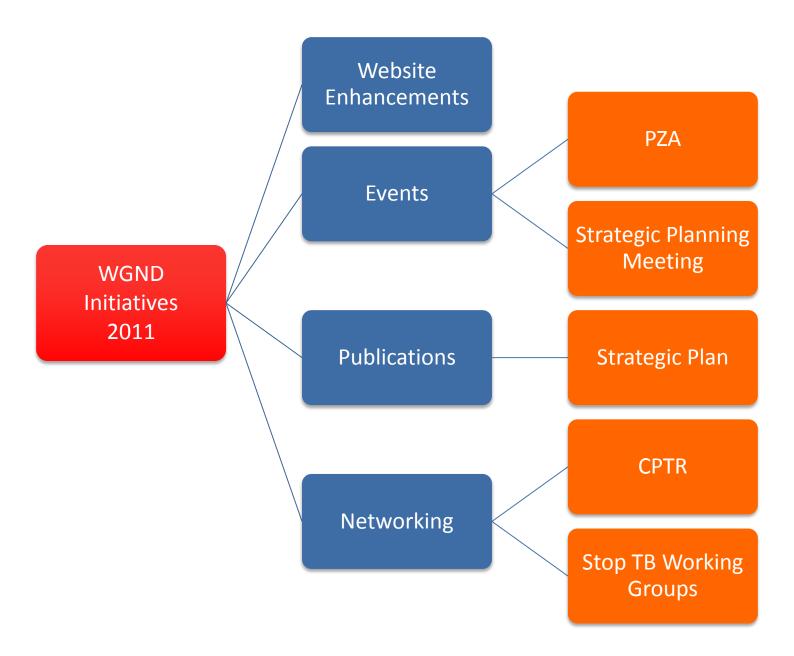




Major Initiatives

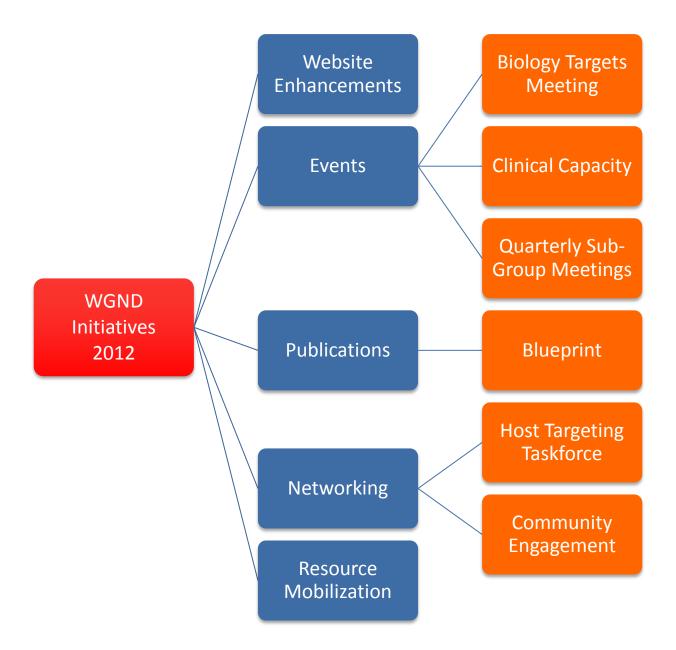










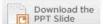




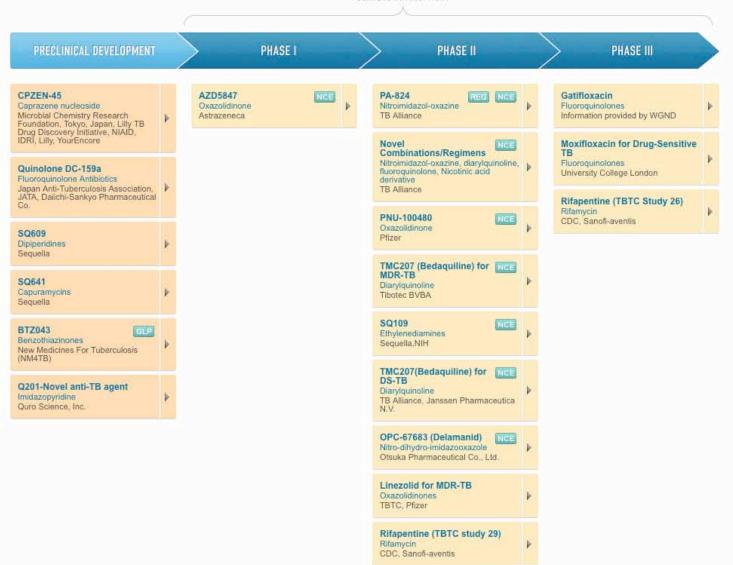


PIPELINE





CLINICAL DEVELOPMENT







DISCOVERY PIPELINE



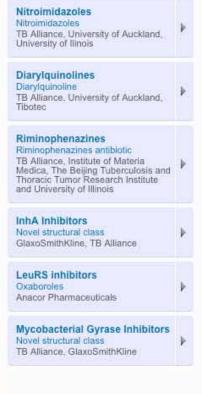


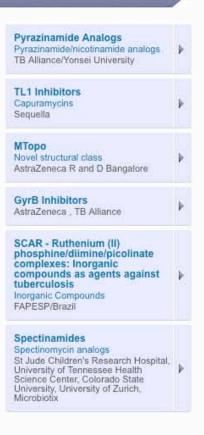
HIT-TO-LEAD

LEAD-OPTIMIZATION

Phenotypic Hit-to-Lead University of Illinois, TB Alliance	Þ
M. tuberculosis Protein Kinase Inhibitors Several chemical scaffolds with various PknA, PknB and PknG selectivity profiles Vertex Pharmaceuticals, Incorporated	Þ
Fungal metabolites Mycosynthetix, University of Illinois at Chicago	Þ
Actinomycete metabolites University of Illinois at Chicago, Myongji University	Þ
DNA metabolism Novel structural class AstraZeneca R and D Bangalore	Þ
Phenotypic hit to lead Multiple novel structural classes Lilly TB Drug Discovery Initiative	Þ
Novel hit-lead programs Novel structural class Lilly TB Drug Discovery Initative	Þ
Phenotype Hit-to-Lead Novel structural classes AstraZeneca R and D Bangalore	Þ

Compounds Shaw Environmental and University of Illinois at Chicago	Þ
Folate Biosynthesis Inhibitors AstraZeneca, TB Alliance	Þ
Phenotypic Hit-to-Lead GlaxoSmithKline, TB Alliance	Þ
Malate Synthase Inhibitors GlaxoSmithKline , Texas A&M University, TB Alliance	Þ
Menaquinone Synthase Inhibitors Colorado State University, TB Alliance	Þ
Inhibitors of Mycobacterium Tuberculosis Energy Metabolism Various Classes UPenn and TB Alliance	Þ
Inhibitors of isoprenoid biosynthesis Lilly TB Drug Discovery Initiative	Þ
Protein Splicing Inhibitors Boston Biomedical Research Institute	Þ











Please Join the conversation



