

What is ISAP ?



- **I** nternational
- **S** ociety for
- **A** ntiinfective
- **P** harmacology

Why did ISAP started ?

Our thinking in the late 80's:

- anti-infective drug usage has long been irrational or not scientifically based on a pharmacodynamic point of view
 - search for low doses for fear of toxicity
 - “errors” in drug dosages at registration
 - misunderstanding of “optimal schedules”
- pharmacokinetics was mostly used to establish “drug presence” rather than to make correlation with efficacy



pharmacodynamics of antiinfective drugs was largely “terra ingognita”

How did it start ?

population
pharmacokinetics

tissue
concentrations

efficacy/toxicity
ratios

AUIC and
fluoroquinolones

postantibiotic
effect
en β -lactam
infusion

once-daily
dosing of
aminoglycosides



Stockholm,
1989 ...

The story of ISAP...

- **thought about in 1989 in Stockholm, Sweden**
- **various names suggested (AIDA, OTELLO, ...) but eventually ISAP chosen in Atlanta, GA, in 1990**
- **officially founded in 1991 in Berlin, Germany**
- **incorporated in 1993 (in South Wales, Australia; now in CA, USA)**
- **108 members worldwide (40% US, 40% Europe, 20% elsewhere)**
 - academy
 - industry

ISAP scientific activities ...

Over the last 10 years, a series of key concepts have emerged from our discussions and work:

- **dose-effect relationships are not the same for all anti-infectives**
 - **beta-lactams vs aminoglycosides, e.g.**
- **some anti-infectives are time-dependent, others not**
- **subinhibitory effects, post-antibiotic effects, cooperation with host defenses modulate the activity of antiinfective drugs**
- **integration of PK/PD allows to fairly predict success or failure, and even emergence of resistance**

ISAP in action in science and clinics

Some achievements:

- **once-daily dosing of aminoglycosides registration or reregistration in several countries**
 - **amikacin, netilmicin (from bid to qd)**
 - **isepamicin (registered essentially for qd dosing)**
- **AUIC / MIC and Cmax / MIC ratios used as guides for phase II / III trials, for treatment optimization and for registration of fluoroquinolones**
 - **trovafloxacin**
 - **moxifloxacin**
- **beta-lactam continuous infusion brought to clinical trials**

ISAP in action with the Regulatory

- FDA (CDER) workshop in 1999
- EMEA (CPMP) Discussion meeting in 1999

ISAP in action for scientific meetings

- **8 independent symposia**
- **8 co-sponsored symposia**
(ICAAC, ECCMID, ICC, ASM General Meeting, ...)

ISAP in action in education

- **4 educational workshops**
- **A WEB site with**
 - **announcements**
 - **slides of main meetings and workshops**

<http://www.isap.org>

Who is ISAP (in 2001) ?

- **President:** George L. Drusano (NY, 2000-2002)
- **Président-elect:** Johan W. Mouton (The Netherlands)
- **Past-presidents:**
 - W.A. Craig (WI; founding president (1991-1994))
 - O. Cars (Sweden; 1994-1996)
 - M.N. Duley (CA; 1996-1998)
 - P.M. Tulkens (Belgium; 1998-2000)
- **Secretary:** J. Mouton (The Netherlands)
- **Treasurer:** J. Scott (CA)

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