Target Concentration Intervention
Dose Individualization using Monitoring of Patient Response
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Objectives

1) Appreciate how a target concentration (TC) strategy is essential for rational clinical use of medicines
2) Understand when and why individual patient monitoring can be used for dose individualization
3) Distinguish target concentration intervention (TCI) from therapeutic drug monitoring (TDM)

Target Concentration in Clinical Use of Medicines

The target concentration approach links pharmacokinetics (PK) with pharmacodynamics (PD) to predict the right dose for a patient.

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How to Find the Target?

Theophylline Target Concentration in Severe Airways Obstruction – 10 or 20 mg/L?
A Randomized Concentration-Controlled Trial

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- Randomized concentration controlled trials are the gold standard

Why does PKPD vary?

- Systematic (predictable)
  - Body size
  - Disease state (liver, kidney)
  - Genotype
  - etc...

- Random (not predictable)
  - Between Subject Variability
  - Within Subject Variability

Predictable Variability
Size and Maturation

¾ Allometry Alone explains 67% of CL variability

¾ Allometry + Maturation explains 80% of CL variability

Differences Remain Even After Accounting for Obvious Features

Figure originally drawn by Dr. N. Sambol CDDS/UMC1997

It shows the frequency distribution of clearance (expressed per kg body weight) for a medicine eliminated largely by metabolism with the distribution of clearance in specific sub-populations. Despite the use of weight, sex, age, disease, concomitant medications there remains a substantial variability of clearance within each sub-population. This remaining variability is what TCI can reduce. The limiting variability for the benefit of TCI is within subject variability.

Theophylline is metabolized by CYP1A2. This enzyme is induced by polycyclic hydrocarbons in cigarette smoke. Enzyme induction reduces variability because there is a maximum biological limit on the extent of enzyme induction.

Note that the naïve per kg method of scaling leads to an apparent higher clearance in children but in fact the clearance in children is the same as adults when scaled based on allometric theory (Anderson BJ, Holford NHG. Mechanistic basis of using body size and maturation to predict clearance in humans. Drug Metab Pharmacokinet. 2009;24(1):25-36.)

Three Ways to Dose

- **Population**
  - Same dose for everyone
    - The dream dosing method! (often used in adults)

- **Group (Covariate guided)**
  - Same dose for similar group
    - e.g. same weight, CLcr, genotype (usually used for children)

- **Individual**
  - Dose determined by individual response
    - e.g. BP, INR, blood conc

The population dosing method is most commonly used but usually just for convenience. This means that some patients are either under-dosed or over-dosed.

The use of weight or renal function to adjust the dose is recommended for some medicines but probably not used as often as it should be.

CLcr = Creatinine clearance, BP = Blood Pressure, INR = International Normalized Ratio

Individual dosing based on response is widely used when the response is easily measured. But sometimes it is used e.g. with anti-HIV medicines, when the within subject variability is large and predictable individualization is not really possible.

Holford NHG, Buclin TMD. Safe and effective variability - A criterion for dose individualization. Ther Drug Monit 2012; 34: 565-68.
Which Drugs for TCI?

1. Usefulness is hard to measure (drug is working when the clinical outcome is not easily observable)
   - Anti-arrhythmics e.g. lignocaine
   - Anti-convulsants e.g. phenytoin
   - Anti-coagulants e.g. warfarin

2. Big unpredictable variability (after using weight, renal function, etc) and small within subject variability
   - Too much variability means either inadequate beneficial effect or too much adverse effect
   - Observing patient response can predict future dose needs

How?
Target Concentration Strategy

1. Choose Target Concentration
2. Determine V and CL using WT etc.
3. Calculate LD and MDR
4. Measure Response (e.g. INR) Revise Target Conc
5. Measure Conc Revise V and CL
6. Goto Step 3

Target Concentrations

<table>
<thead>
<tr>
<th>Drug</th>
<th>Target Conc</th>
<th>Clearance</th>
<th>Volume of distribution</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aminoglycosides</td>
<td>Peak 20 mg/L</td>
<td>6 L/h</td>
<td>18 L</td>
</tr>
<tr>
<td></td>
<td>Css 3 mg/L</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tacrolimus**</td>
<td>7 mcg/L</td>
<td>20 L/h</td>
<td>100 L</td>
</tr>
<tr>
<td>Phenytoin</td>
<td>10 mg/L</td>
<td>Vmax=415 mg/d, Km=4 mg/L</td>
<td>45 L</td>
</tr>
<tr>
<td>Digoxin</td>
<td>1 ng/mL</td>
<td>9 L/h</td>
<td>500 L</td>
</tr>
<tr>
<td>Theophylline</td>
<td>10 mg/L</td>
<td>3 L/h</td>
<td>35 L</td>
</tr>
</tbody>
</table>

* 24 hour dosing ** whole blood

The first reason for using TCI uses a response, such as blood pressure, as a substitute for being able to measure the clinical disease state that is being treated. When the medicine is working well or it is not working at all the clinical disease state may appear to be the same. It is assumed that trying to reach a typical response that is usually associated with benefit is better than giving everyone the same dose.

The second reason for using TCI is when group based dosing (e.g. using weight) is not enough to reduce the variability so that the medicine can be used safely and effectively. TCI can only work however if the within subject variability (that cannot be influenced by TCI) is small enough so that dose individualization is really predictive for future use of the medicine in the same patient.

Target concentrations and PK parameters are known for most medicines which are helped by TCI.
Target Effects and Pharmacodynamic Parameters for Selected Medicines. Emax is the maximum effect due to the drug, C50 is the concentration producing 50% of Emax. PEFR is peak expiratory flow rate.

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<tr>
<td>Aminoglycosides</td>
<td>&quot;cure&quot;</td>
<td>?</td>
<td>?</td>
</tr>
<tr>
<td>Tacrolimus</td>
<td>&quot;prevention of rejection&quot;</td>
<td>?</td>
<td>?</td>
</tr>
<tr>
<td>Phenytoin</td>
<td>&quot;prevention of seizures&quot;</td>
<td>?</td>
<td>?</td>
</tr>
<tr>
<td>Digoxin</td>
<td>&quot;control of atrial fibrillation&quot;</td>
<td>?</td>
<td>?</td>
</tr>
<tr>
<td>Theophylline</td>
<td>&quot;normal PEFR&quot;</td>
<td>344 L/min</td>
<td>11 mg/L</td>
</tr>
<tr>
<td>Warfarin</td>
<td>INR 2-3</td>
<td>100%*</td>
<td>1.5 mg/L</td>
</tr>
</tbody>
</table>

* Inhibition of prothrombin complex synthesis

The pharmacodynamic parameters for medicines that use TCI are typically not well known. This is a reflection of the difficulty of measuring a clinical response that can be related to concentration.


Determine Group V and CL (predictable variability)

- **Volume of Distribution**
  - size: \( V = V_{pop} \times \frac{WT}{WT_{std}} \)
  - body composition

- **Clearance**
  - size: \( CL = C_{Lpop} \times \left( \frac{WT}{WT_{std}} \right)^{3/4} \)
  - renal function
  - hepatic function
  - concomitant drugs

WT=patient weight
WTstd=standard weight e.g. 70 kg
Vpop, CLpop=population volume and clearance in a standard subject e.g. 70 kg.
Calculate LD and MDR
e.g. Gentamicin

\[ \text{LD} = \text{TC} \times V = 20 \text{ mg/L} \times 20 \text{ L} = 400 \text{ mg} \]

\[ \text{MDR} = \text{TC} \times \text{CL} = 3 \text{ mg/L} \times 6 \text{ L/h} = 18 \text{ mg/h} \]
\[ = 400 \text{ mg/day} \]

Number of Samples
- Most medicines 1
- Gentamicin 2

Timing of Sample
- As soon as possible
  - Most medicines Middle of dosing interval
  - Gentamicin "peak" and "trough"

When to Measure Concs?
Goal is to estimate PK e.g. CL

A concentration in the middle of the dosing interval \( (C_{\text{tmid}}) \) will be closer to the average steady state concentration \( (C_{\text{ss}}) \) then either a peak or trough concentration.

Clearance is easily calculated from \( CL = \frac{\text{Dose Rate}}{C_{\text{ss}}} \) which can be approximated by \( CL = \frac{\text{Dose Rate}}{C_{\text{tmid}}} \).

A rational approach to measuring drug concentrations is based on using the measurement to predict pharmacokinetic parameters – most commonly clearance.

The least informative time to measure concentrations is just before the next dose (the ‘trough’ concentration) unless this is paired with another ‘peak’ concentration. This is because clearance determines the average concentration. So measuring a concentration in the middle of the dosing interval will be closer to the average and therefore more useful for predicting clearance.

Gentamicin concentrations vary widely in a dosing interval so two concentrations are needed to reliably estimate clearance. The sooner the sample is taken the sooner concentrations can be used to improve dosing.
Gentamicin concentrations with once a day dosing vary considerably. The trough concentration at 24 h is often unmeasurable because it is below the limit of quantitation. Concentrations are best measured 1 h and 8 h after the dose.

In this example a dose of 240 mg was given every 24 hours. V can be approximated from the first conc (C1 = 13 mg/L) assuming about 15% of the dose is eliminated in 1 hour:

\[ V = \frac{0.85 \times \text{Dose}}{C_1} = \frac{0.85 \times 240 \text{ mg}}{13 \text{ mg/L}} = 15.7 \text{ L} \]

Half-life can be estimated from C1 at time T1 (1 h) and the second conc (C2 = 2.5 mg/L) at time T2 (8 h):

\[ k = \ln\left(\frac{C_1}{C_2}\right)/(T_2 - T_1) = \ln\left(\frac{13}{8}\right)/(8 - 1) = 0.236 \text{ h}^{-1} \]

\[ \text{Thalf} = \frac{\ln(2)}{K} = 2.94 \text{ h} \]

\[ CL = V \times k = 3.7 \text{ L/h} \]

TDM or TCI?

- **Therapeutic Drug Monitoring**
  - TDM Therapeutic Range
  - Sub-optimal at borders of the range

- **Target Concentration Intervention**
  - TCI Single Target
  - Accurate

Target concentration intervention is a science based method that uses pharmacokinetic and pharmacodynamic principles to identify how patients are different and uses PK guided dose individualization to achieve a precise therapeutic target. It has been shown to improve clinical outcome as well as being a cost-effective use of health resources.


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