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Size does matter : drug glucuronidation in children

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Section IV

Paediatric Model Evaluation

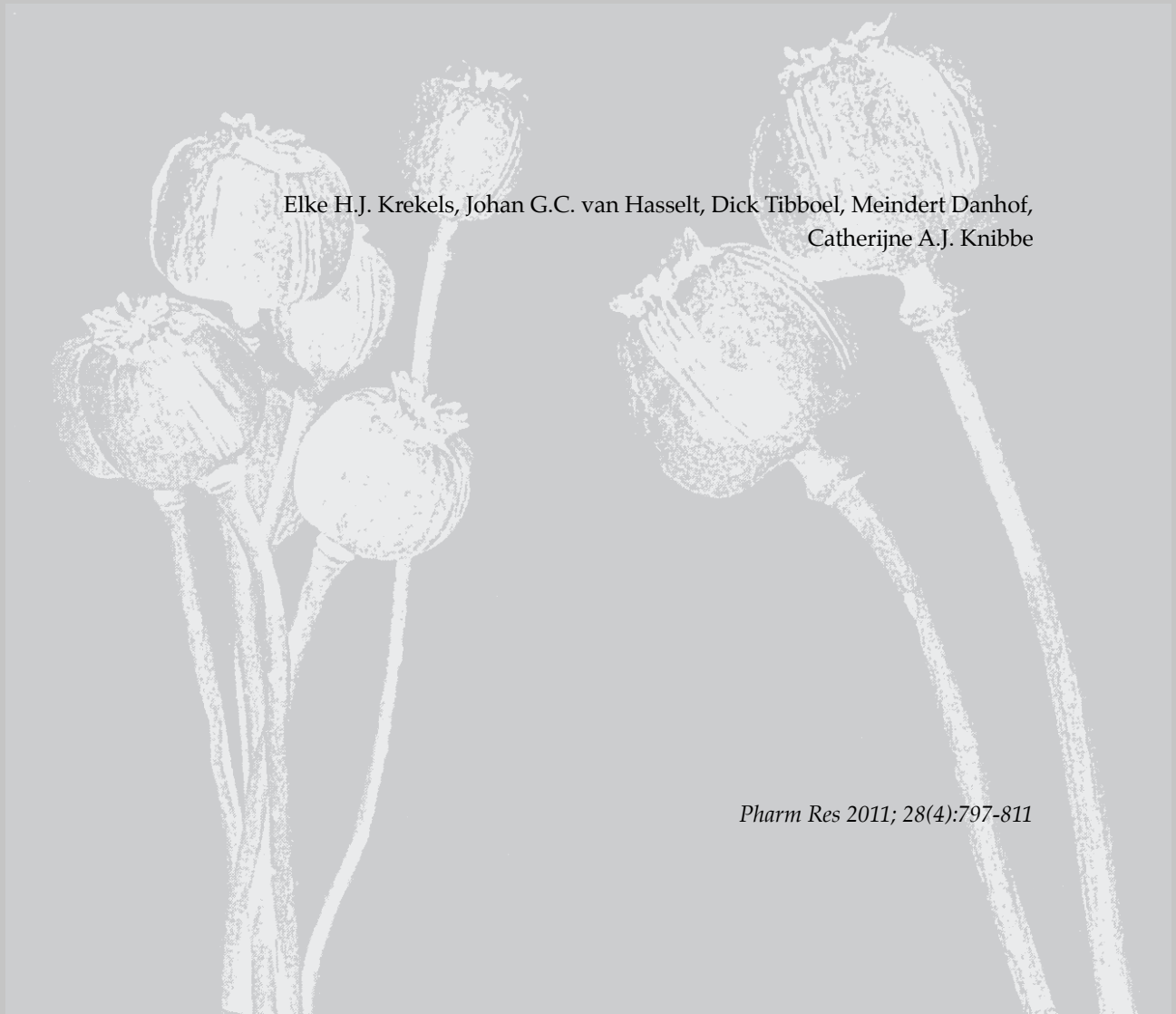


Chapter 8

Systematic Evaluation of the Descriptive and Predictive Performance of Paediatric Morphine Population Models

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Abstract

Purpose: A framework for the evaluation of paediatric population models is proposed and applied to two different paediatric population pharmacokinetic models for morphine. One covariate model was based on a systematic covariate analysis, the other on fixed allometric scaling principles.

Methods: The six evaluation criteria in the framework were 1) number of parameters and condition number, 2) numerical diagnostics, 3) prediction-based diagnostics, 4) η -shrinkage, 5) simulation-based diagnostics, 6) diagnostics of individual and population parameter estimates *versus* covariates, including measurements of bias and precision of the population values compared to the observed individual values. The framework entails both an internal and external model evaluation procedure.

Results: The application of the framework to the two models resulted in the detection of over-parameterization and misleading diagnostics based on individual predictions caused by high shrinkage. The diagnostic of individual and population parameter estimates *versus* covariates proved to be highly informative in assessing obtained covariate relationships. Based on the framework, the systematic covariate model proved to be superior over the fixed allometric model in terms of predictive performance.

Conclusions: The proposed framework is suitable for the evaluation of paediatric (covariate) models and should be applied to corroborate the descriptive and predictive properties of these models.

8.1 Introduction

Whereas many diagnostic and validation tools are available for the evaluation of population models in the adult population, these tools may not always directly suffice in the paediatric population due to the heterogeneity of this special population and the scarcity of the datasets. In this study a framework including six evaluation criteria is presented for the systematic assessment of the descriptive and predictive properties of paediatric (covariate) models that takes these specific issues into consideration.

In paediatric population pharmacokinetic (PK) models, the influence of the many physiological changes that take place in the paediatric age-range are reflected in covariate relationships that are usually based on bodyweight and/or age. However, since bodyweight and age are naturally correlated in the paediatric population there is a debate on how to incorporate the influence of the physiological changes in paediatric population PK models. Bodyweight and age can either be regarded as regular covariates whose predictive properties on PK parameters are evaluated together with other covariates in a systematic covariate analysis by formally testing them for significance and only retaining them in the model if they statistically improve the model fit (Chapter 3) ^[1,2]. Alternatively, bodyweight can be included *a priori* into paediatric PK models by the use of a bodyweight-based allometric equation with a fixed exponent of 0.75 for clearance and 1 for distribution volume. These equations can subsequently be augmented by estimated age-based functions of various forms ^[2-4].

In recent years two different population PK models for morphine and its two major pharmacologically active metabolites morphine-3-glucuronide (M3G) and morphine-6-glucuronide (M6G) in children younger than three years have been published (Chapter 3)^[3]. The model by Knibbe *et al.* (Chapter 3) was developed using a systematic covariate analysis. The model by Bouwmeester *et al.* ^[3] was developed using fixed allometric scaling principles in conjunction with estimated age-based functions. As these models were developed using similar datasets, they provide both an example for the assessment of the developed framework for the evaluation of paediatric (covariate) models, as well as an opportunity to directly compare the performance of these two fundamentally different paediatric covariate models.

8.2 Materials and Methods

8.2.1 Models and Data

Figure 1 shows a schematic representation of the two models that are evaluated in the current analysis. In the model by Knibbe *et al.* (Chapter 3) the maturation of the formation and elimination clearances of the morphine glucuronides was found to be best described by a bodyweight-based exponential equation with an estimated exponent of 1.44. Within this exponential equation the formation clearance of the glucuronides was found to be significantly reduced in neonates younger than ten days. Distribution volumes were estimated to scale linearly with bodyweight. This model will be referred to as the systematic covariate model. In the model by Bouwmeester *et al.* [3] bodyweight was included *a priori* using an allometric equation with fixed exponents of 0.75 for clearance and 1 for distribution volume. Three exponential equations based on postnatal age (PNA) augmented the model, one equation for distribution volumes, one for the formation of the morphine metabolites and one for the elimination of the metabolites. Bilirubin concentration and creatinine concentrations were also incorporated to the model as covariates for the formation and elimination of the glucuronides respectively. This model will be referred to as the fixed allometric model.

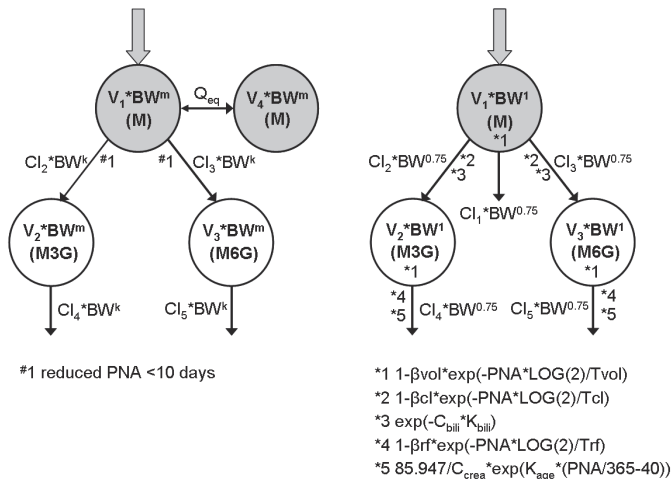


Figure 1. Schematic representation of the systematic covariate model (Chapter 3) (A) and the fixed allometric model [3] (B). M = morphine, M3G = morphine-3-glucuronide, M6G = morphine-6-glucuronide, V = distribution volume of the designated compartment, Cl = clearance of designated route, Q = intercompartmental clearance, PNA = postnatal age, k and m = exponential scaling constants, β = fraction below adult values at birth and T = maturation half-life for distribution volume (vol), formation clearance of the metabolites (cl), and elimination clearance of the metabolites (rf), C = plasma concentration and K = scaling constant for bilirubin (bili) and creatinine (crea).

The systematic covariate model (Knibbe *et al.* (Chapter 3)) was developed using two datasets ^[5,6], while the fixed allometric model (Bouwmeester *et al.* ^[3]) was developed using only one of these two datasets ^[5]. To allow for a direct comparison, the systematic covariate model was refit with the data from the one common dataset. This dataset will be referred to as the internal dataset of this study and consists of postoperative term neonates, infants and children up to the age of three years on a continuous or intermitted intravenous morphine regimen ^[5].

For the external evaluation of the two paediatric covariate models in the current analysis, five previously published datasets ^[6-10] were used. These external datasets included the same patient population as the internal dataset with the exception that their PNA ranged up to only one year instead of three years. In addition, two external datasets included preterm neonates ^{[6][7]}, which is a younger age-range than the age-range in the internal dataset ^[5] used for model building. An overview of all datasets is given in table I.

Table I. Overview of the internal dataset (Int.1) and the external datasets (Ext.1-5) used for model building and external model evaluation.

Dataset	Patient Population	Number of Patients	Postnatal Age in days (median, IQR)	Bodyweight in g (median, IQR)
Int.1. ^[5]	Post-operative term neonates, infants and children.	183	97 (8 – 286)	4700 (3100 – 8000)
Ext.1. ^[8]	Post-operative term neonates and infants	28	14 (0 – 70)	3100 (2550 – 4000)
Ext.2. ^[9]	Post-operative term neonates and infants	9	10.5 (3 – 135)	3800 (3000 - 5000)
Ext.3. ^[10]	Term neonates and infants on artificial ventilation	12	13 (6 – 80)	3050 (2675 – 6900)
Ext.4. ^[6]	Preterm and term neonates on artificial ventilation	63	0.4 (0.2 – 0.5)	1180 (862.5 – 1760)
Ext.5. ^[7]	Preterm neonates on artificial ventilation	41	1 (1 – 2)	1035 (892.5 – 1295)

Int. = internal dataset, Ext. = external dataset, M = morphine, M3G = morphine-3-glucuronide, M6G = morphine-6-glucuronide

8.2.2 Model Evaluation

All model fitting and model-based simulations in the current study were performed using NONMEM VI (ICON, Ellicott City, MD).

The framework for the evaluation of paediatric population models is composed of the following six evaluation criteria and tools:

- 1) total number of parameters and condition number of the model. The latter was obtained by taking the ratio of the largest and smallest eigenvalue of the covariance matrix of the estimate from the NONMEM output.
- 2) numerical diagnostics by means of a bootstrap analysis using the PsN software package ^[11]. One hundred datasets were resampled with replacement from the internal dataset and refit to the models. The parameter estimates for fixed and random effects obtained in every separate run were summarized in terms of mean and relative standard errors (RSE) for each parameter. Runs that did not minimize successfully were excluded from the analysis.
- 3) prediction-based diagnostics by means of basic goodness-of-fit plots. Both the individual and population predicted concentrations were plotted *versus* the concentrations that were actually observed in these datasets. Plots were made for both the internal and external datasets and for the population as a whole as well as for stratified subsets based on age (0-1 month, 1 month – 1 year, 1-3 years). Mirror plots were created to serve as a reference for these predicted *versus* observed plots ^[12].
- 4) η -shrinkage as defined by Karlsson and Savic. ^[12], which was calculated for all model parameters for which inter-individual variability was estimated.
- 5) simulation-based diagnostics by means of normalized prediction distribution errors (NPDE) ^[13]. Both the internal and merged external datasets were simulated 1000 times with inclusion of the inter-individual variability and residual error. Using the NPDE add-on package for R (version 1.2) ^[14] a cumulative distribution was assembled for each observation with the 1000 simulated concentrations and subsequently the value of the cumulative distribution at the observed concentration was determined. An inverse function of the normal cumulative density function was then applied to these data to obtain what are called the normalize prediction distribution errors. The NPDEs are presented in a total distribution, *versus* time and *versus* the concentration. This analysis was also performed on the population as a whole and on the stratified subsets described in item 3.
- 6) Individual and population parameter estimates *versus* the most predictive covariate in the model. In both models, bodyweight was the most predictive covariate. Total morphine clearance was defined as the sum of Cl_1 and Cl_2 for the systematic covariate model (see figure 1A) and of Cl_v , Cl_1 , and Cl_2 for the fixed allometric model (see figure 1B). The elimination clearances of the metabolites (Cl_3 and Cl_4) and the distribution volume of the central morphine compartment (V_1) were directly compared between the two models. To numerically quantify the bias and precision of the model predicted parameter values compared to the observed parameter values in the internal datasets, Mean Prediction Error (MPE,

equation 1) and the Root Mean Square Error (RMSE, equation 2) were calculated respectively for both models.

$$MPE = \frac{\sum \frac{(predicted - observed) * 100}{observed}}{n} \quad (\text{equation 1})$$

$$RMSE = \sqrt{\frac{\sum (predicted - observed)^2}{n}} \quad (\text{equation 2})$$

In these equations predicted parameter values were the population predicted values for each individual by both models and the observed parameter values were the individually observed *post hoc* parameter values for that individual. MPE and RMSE were calculated separately for the strata described before. A table with MPE and RMSE was also constructed for the external datasets. This table was also stratified based on age with the following strata: preterm neonates (PNA < 1 month and postmenstrual age (PMA) at birth < 36 weeks), term neonates (PNA < 1 month and PMA at birth ≥ 36 weeks), toddlers (PNA 1 month – 1 year).

8.3 Results

Table II A and B lists the parameter estimates as obtained with the fit of the internal dataset by the systematic covariate model and the fixed allometric model, respectively. With a total of 18 model parameters, the systematic covariate model described the fixed and random effects with fewer parameters than the fixed allometric model, which contains 35 model parameters. The condition number of the systematic covariate model was 293, which is well below the critical value for the indication of serious ill-conditioning of 1000 [15]. For the fixed allometric model the condition number was 10698, which is more than ten-fold higher than the critical value.

Table II. Parameter estimates from the model fit, number of model parameters and parameter estimates from the bootstrap procedure for the systematic covariate model (A) and for the fixed allometric model (B) obtained with the internal dataset. Parameter names are explained in figure 1.

A.

Parameters	<i>Model fit</i>	<i>Bootstrap</i>
	Value (RSE%)	(98 out of 100 successful) Mean value (RSE%)
Systematic covariate model		
Fixed effects (n=10)		
k = exponential scaling factor on clearance	1.49 (3.6)	1.49 (3.42)
m = exponential scaling factor on distribution volume	1 fixed	1 fixed
Cl _{1 PNA < 10d} (ml/min/kg ^k)	3.68 (9.0)	3.68 (8.15)
Cl _{1 PNA > 10d} (ml/min/kg ^k)	8.04 (11.0)	8.01 (10.1)
Cl _{2 PNA < 10d} (ml/min/kg ^k)	0.423 (11.1)	0.42 (14.4)
Cl _{2 PNA > 10d} (ml/min/kg ^k)	0.623 (10.2)	0.62 (13.0)
Cl ₃ (ml/min/kg ^k)	1.84 (9.9)	1.85 (9.73)
Cl ₄ (ml/min/kg ^k)	0.955 (9.0)	0.95 (12.6)
Q _{eq} (ml/min)	40.8 (24.1)	43.8 (42.7)
V ₁ = V ₄ (l/kg)	1.64 (8.2)	1.65 (9.44)
V ₂ = V ₃ (fraction of V ₁)	0.157 (21.2)	0.161 (27.0)
Inter-individual variability (n=5)		
ω ² Cl ₁	0.0809 (24.0)	0.0774 (22.9)
ω ² Cl ₃	0.256 (27.1)	0.263 (29.1)
ω ² Cl ₄	0.110 (15.2)	0.110 (15.5)
ω ² Cl ₃ -Cl ₄ covariance	0.128 (18.2)	0.126 (19.8)
ω ² V ₁	0.162 (17.9)	0.168 (20.7)
Residual variance (n=3)		
σ ^{2, prop} (morphine)	0.440 (14.7)	0.431 (14.0)
σ ^{2, prop} (M3G)	0.261 (27.2)	0.243 (26.0)
σ ^{2, prop} (M6G)	0.0894 (15.7)	0.0894 (16.4)

B.

<i>Parameters</i>	<i>Model fit</i>	<i>Bootstrap</i> (46 out of 100 successful)
	<i>Value (RSE%)</i>	<i>Value (RSE%)</i>
Fixed allometric model		
Fixed effects (n=14)		
Cl_0 (l/h/70kg ^{0.75})	3.12 (117)	2.59 (43.4)
Cl_1 (l/h/70kg ^{0.75})	64.3 (18.0)	55.2 (32.3)
Cl_2 (l/h/70kg ^{0.75})	3.63 (14.0)	3.99 (65.6)
Cl_3 (l/h/70kg ^{0.75})	17.4 (16.0)	7.23 (38.0)
Cl_4 (l/h/70kg ^{0.75})	5.8 (20.2)	5.43 (23.4)
V_1 (l/70kg)	136 (59.3)	147 (34.9)
V_2 (l/70kg)	23 fixed	23 fixed
V_3 (l/70kg)	30 fixed	30 fixed
Bcl	0.834 (6.41)	0.894 (8.35)
Tcl (days)	88.3 (37.4)	65.2 (176)
Brf	0.832 (9.74)	0.814 (10.7)
Trf (days)	129 (49.8)	136 (45.4)
Bvol	0.391 (28.4)	0.388 (38.1)
Tvol (days)	26.3 (72.2)	26.7 (58.9)
K_{age}	0.0141 (140)	0.0201 (36.3)
K_{bili}	-0.00203 (33.2)	-0.00207 (35.3)
Inter-individual variability (n=16)		
$\omega^2 Cl_0$	1.37 (104)	1.80 (69.9)
$\omega^2 Cl_1$	0.346 (20.9)	0.916 (74.9)
$\omega^2 Cl_2$	0.675 (29.3)	1.21 (75.8)
$\omega^2 Cl_3$	0.185 (20.8)	0.764 (41.7)
$\omega^2 Cl_4$	0.545 (32.1)	1.39 (87.6)
$\omega^2 V_1$	0.351 (29.1)	1.54 (121)
full omega block on all eta's except Cl_0	Data not shown	Data not shown
Residual variance (n=5)		
$\sigma^2_{prop morphine}$	0.128 (11.6)	0.503 (105)
$\sigma^2_{add M3G}$ (ng/ml)	50.3 (36.2)	136 (217)
$\sigma^2_{prop M3G}$	0.118 (27.0)	2.21 (80.0)
$\sigma^2_{add M6G}$ (ng/ml)	0.198 (26.0)	1.31 (148)
$\sigma^2_{prop M6G}$	0.0925 (16.9)	0.249 (53.0)

In tables II A and B the parameter estimates obtained with the bootstrap analyses are presented as well. The parameter estimates of the bootstrap deviated more from the values obtained in the initial model fit for the fixed allometric model compared to the systematic covariate model. For both the model fit and the bootstrap procedure with the fixed allometric model, the overall precision of the parameter estimates was lower than the systematic covariate model as expressed by the higher relative standard error (RSE) of the parameter estimates. Additionally, for the bootstrap, using the fixed allometric model only 46 out of 100 model refits minimized successfully, whereas 98 out of 100 model refits successfully minimized using the systematic covariate model.

Figure 2 shows the individual predicted concentrations *versus* observed concentrations for morphine and its metabolites as obtained with the internal dataset for both the systematic covariate model (A) and the fixed allometric model (B). This figure shows a slightly better description of individual concentrations by the fixed allometric model compared to the systematic covariate model, especially for the mother compound morphine.

Figure 3 shows the population predicted concentrations *versus* observed concentrations obtained with the internal as well as the five external datasets. This figure shows the systematic covariate model to be superior over the fixed allometric model in the predictions of population concentrations in the datasets. The predictions for the systematic covariate model are only slightly biased and since this bias is also observed in the mirror plots (data not shown) this is not indicative of model misspecification. For the fixed allometric model on the other hand significant bias towards under-prediction can be observed, that did not correspond to trends observed in the mirror plots (data not shown).

For both models, stratification of the plots of the predicted *versus* observed concentrations into the different age-groups showed no differences in model performance (data not shown).

Table III shows the percentage of η -shrinkage for the parameters for which inter-individual variability was identified in each of the two models. Both models have parameters for which shrinkage is relatively high (>20%), indicating that the individual data in the internal dataset is not rich in information about these parameters.

Table III. Percentage η -shrinkage in both models for the parameters for which inter-individual variability was identified.

	Cl ₀	Cl ₁	Cl ₂	Cl ₃	Cl ₄	V ₁
Systematic covariate model (%)	-	29.6	-	8.26	5.76	30.3
Fixed allometric model (%)	52.3	10.8	13.2	21.0	18.9	17.7

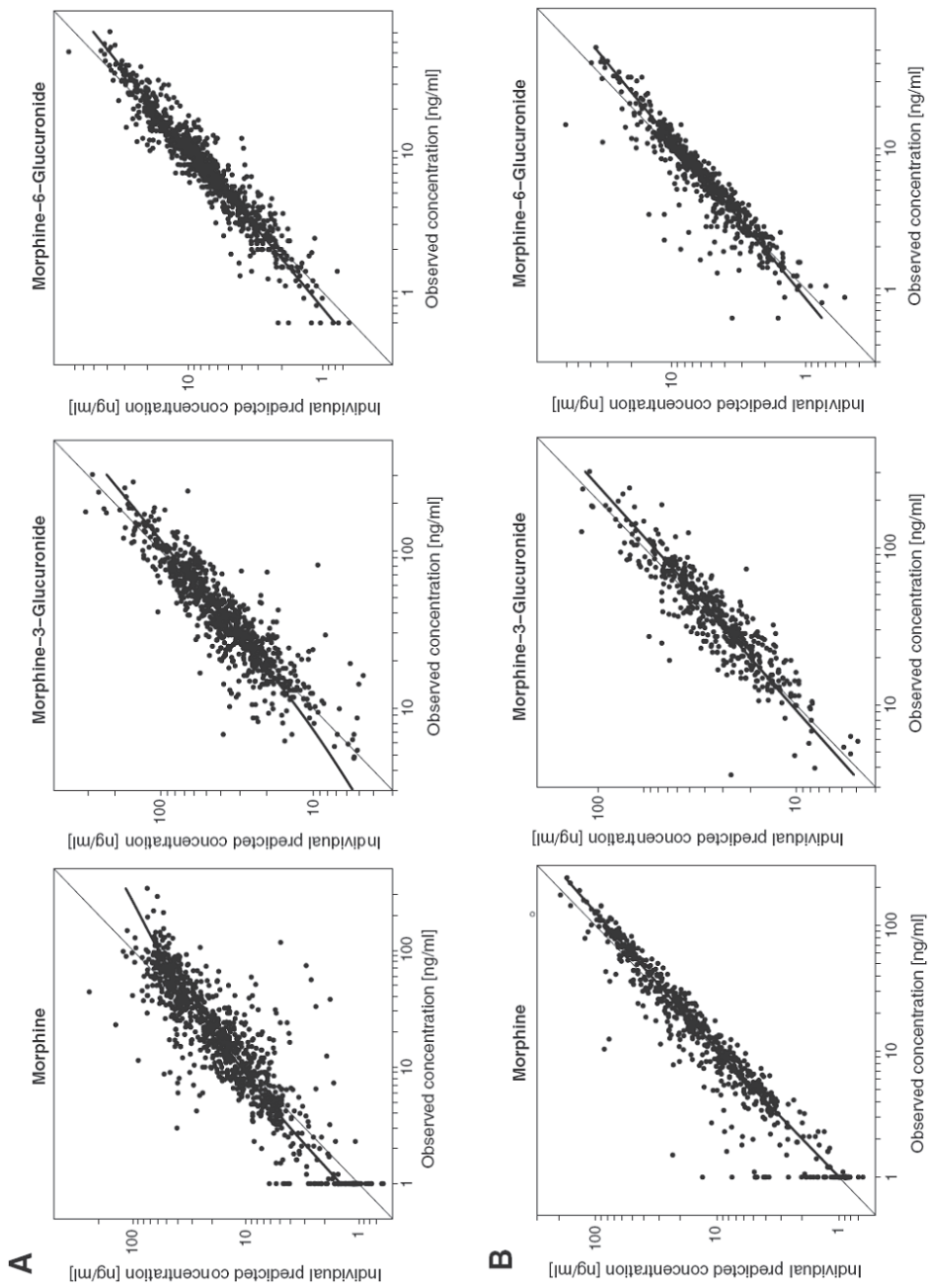


Figure 2. Individual predicted concentrations versus observed concentrations including a loess curve of morphine and its metabolites in the internal dataset. Predictions by the systematic covariate model are depicted in panel A and predictions by the fixed allometric model in panel B.

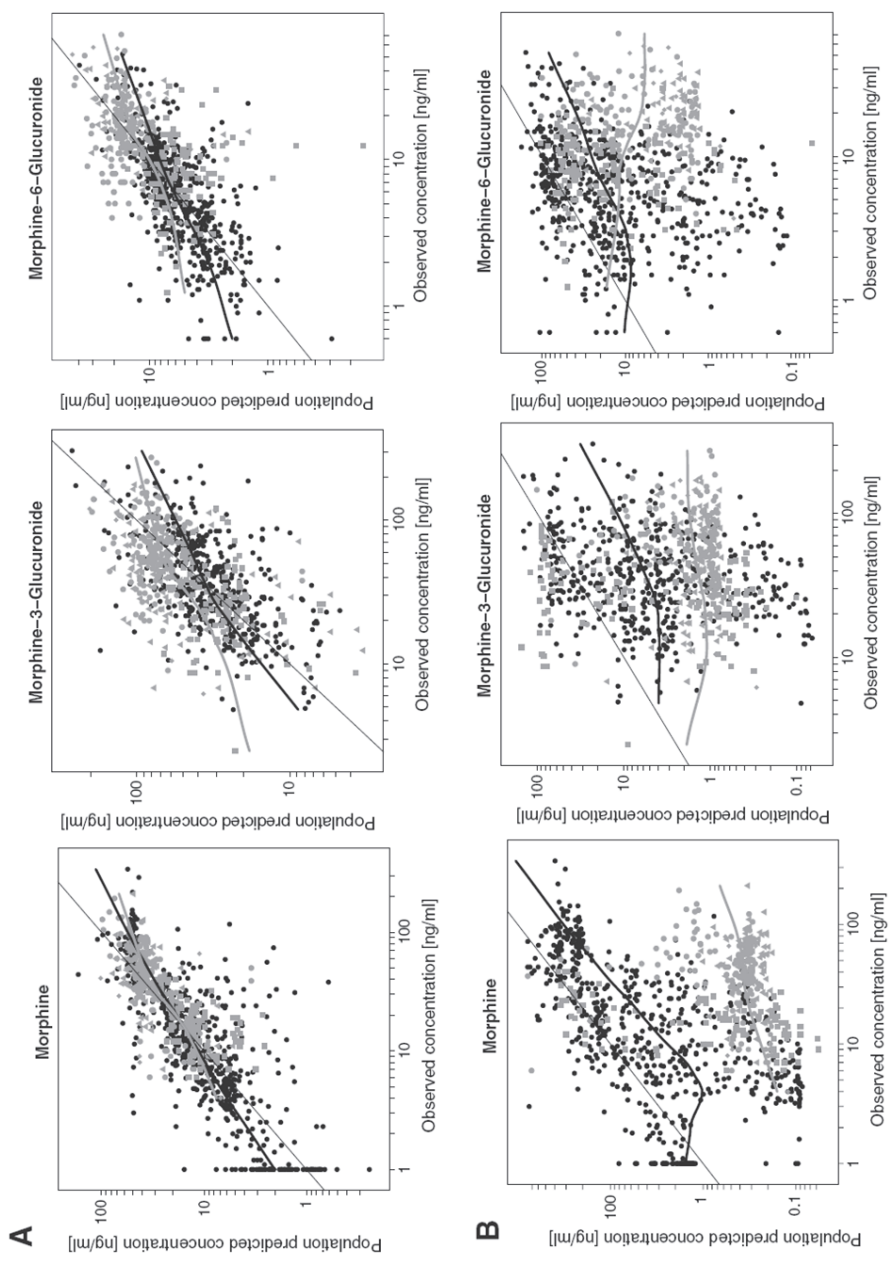


Figure 3. Population predicted concentrations versus observed concentrations including loess curves of morphine and its metabolites for the systematic covariate model (A) the fixed allometric model (B). Data points in black originate from the internal dataset and data points in grey from the external datasets. Different symbols are used for different external datasets: ■ = Ext. 1 ^[8], ● = Ext 2 ^[9], ◆ = Ext 3 ^[10], * = Ext 4 ^[6], ▲ = Ext 5 ^[7].

For both models the results of the NPDE analysis with the internal dataset are depicted in figure 4. For the systematic covariate model no trends in time or concentration are observed and the mean of the distribution of NPDEs is close to 0 while the variance is slightly lower than 1. The trends observed in the plots for the fixed allometric model are indicative of an under-prediction, which appears to be relatively constant over time but to increase with decreasing concentrations.

The results of the NPDE analysis with the external datasets confirmed the results obtained with the internal dataset for both models (data not shown). Stratification into different age-groups also revealed similar results for the three age-groups for each of the models (data not shown).

The plots of the individual *post hoc* parameter estimates and population predicted parameter estimates for total morphine clearance, the clearances of the metabolites and distribution volume of the central morphine compartment *versus* bodyweight for both models are shown in figure 5. For the systematic covariate model, total morphine clearance is composed of Cl_1 and Cl_2 (see figure 1A) which both have different population values for children older and younger than ten days resulting in two different lines of population parameter estimates. For the fixed allometric model, total morphine clearance is composed of Cl_r , Cl_1 and Cl_2 (see figure 1B). The larger number of additional covariates (age, and bilirubin and creatinine concentration) on the structural parameters in the fixed allometric model results in scattered lines for the population parameter estimates of this model. This figure shows that for the systematic covariate model the population predicted values describe the individual *post hoc* values without bias for all parameters, whereas for the fixed allometric model the population predicted values are biased compared to the individual *post hoc* values for all parameters.

Table IV numerically quantifies the bias (MPE) in the population predicted parameter values compared to the individually observed parameter values for both models. The RMSE in this table quantifies the precision of the population prediction. It can be seen that for the systematic covariate model, mean bias in the population predictions stays well below 25% and remains relatively constant over the age-ranges for all parameters. For the fixed allometric model, bias in the population predictions reaches up to 250% and especially in the clearance of the metabolites an increasing trend towards over-prediction is observed with increasing age.

Table V shows the mean bias and precision in the predictions of the model parameters in the external dataset stratified in three age-groups. As can be expected, for both models the bias in the parameter predictions of the external datasets is generally larger than for the internal dataset, however for the systematic covariate model it still remains below 35.6%, whereas for the systematic covariate model the values are between 25% and 300%.

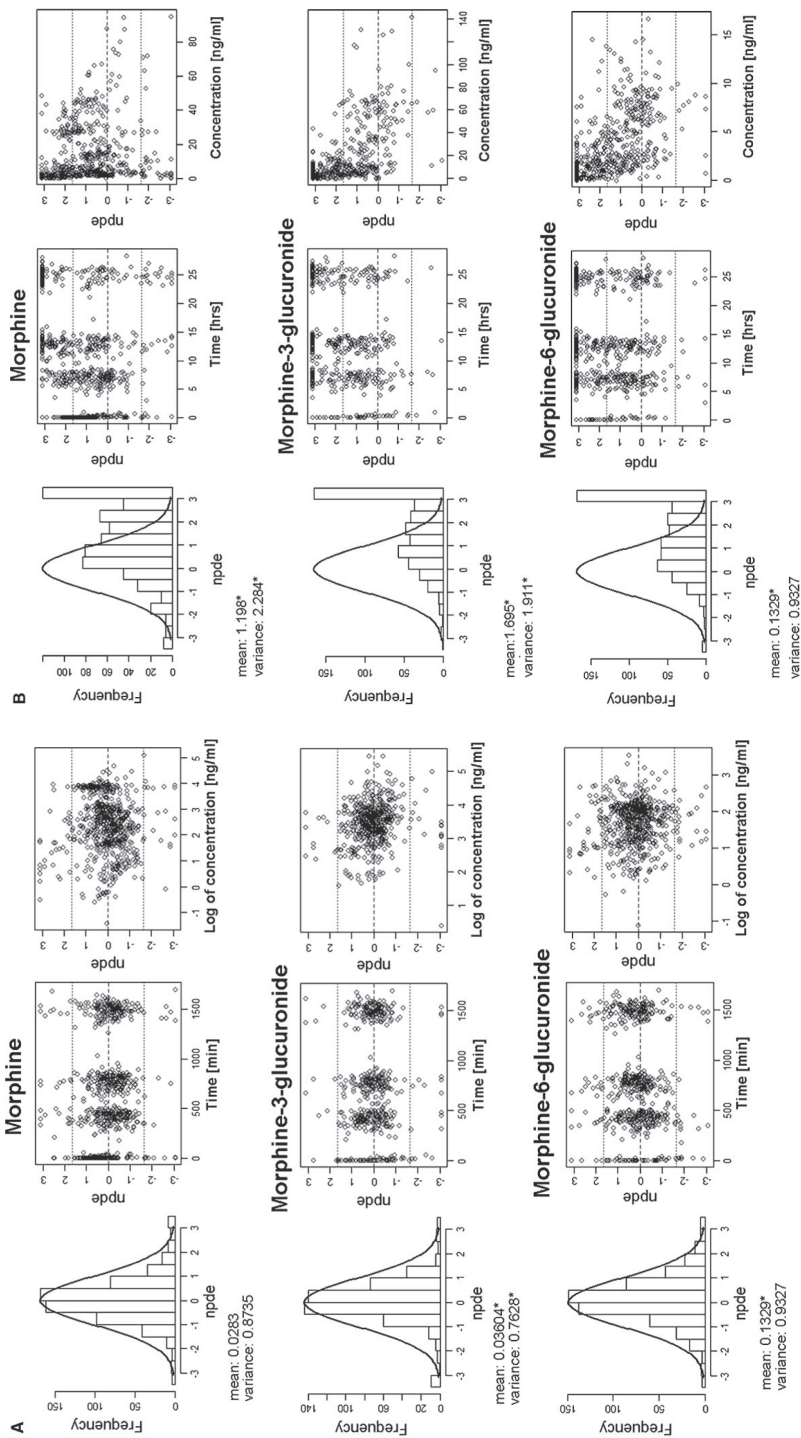


Figure 4. Result of the NPDE analysis for morphine and its metabolites using the internal dataset with the systematic covariate model (A) and the fixed allometric model (B). In the histograms the distributions of the NPDEs for morphine and its metabolites in the total dataset are shown. The solid line depicts a normal distribution and the values below specify the mean and standard deviation of the observed NPDE distribution in the histogram. A statistically significant deviation of the distribution from a mean of 0 and a standard deviation of 1 is indicated with an asterisk (*). The distributions of NPDEs in time after first dose and against the observed concentrations are also shown. As for the systematic covariate model log-transformed data have been used, the last plot shows the NPDE against the log-value of the observed concentration.

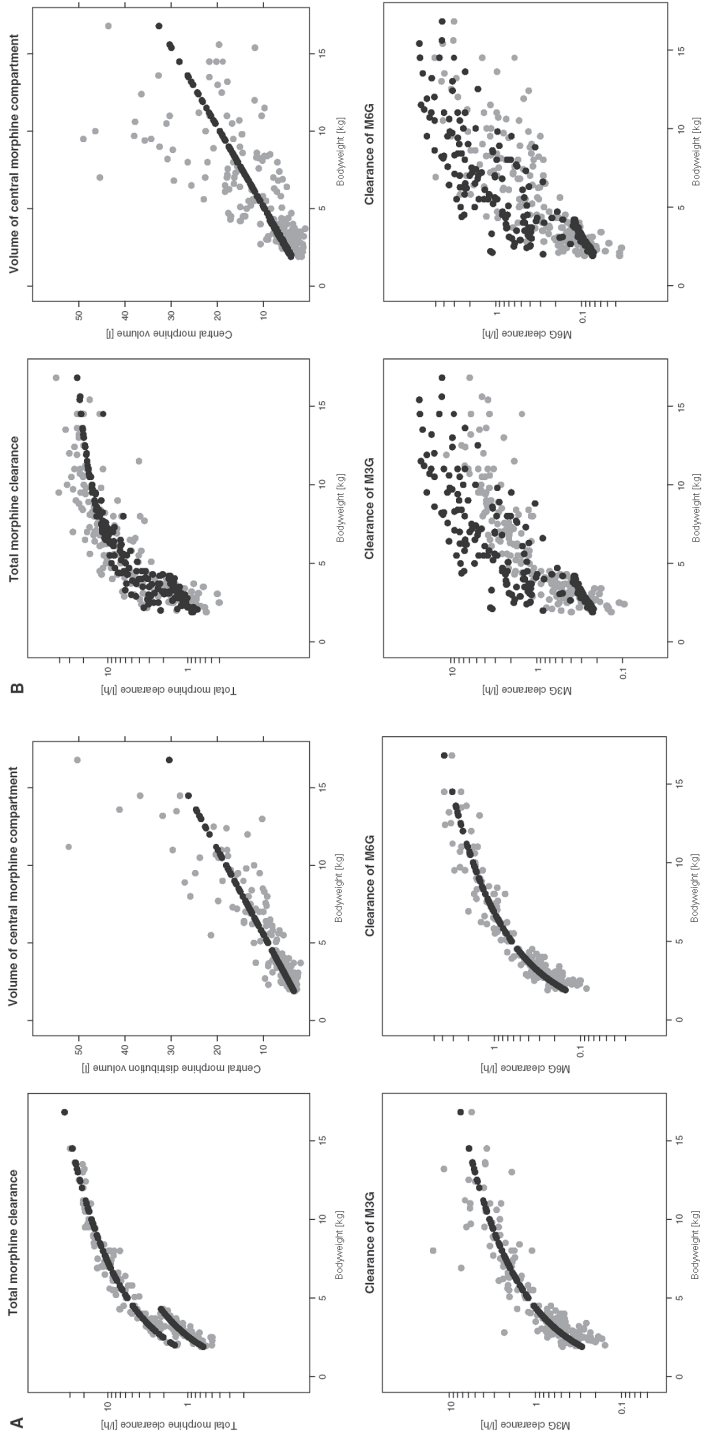


Figure 5. Individual post hoc parameter estimates (grey) and population predicted parameter estimates (black) for total morphine clearance ($Cl_0+Cl_1+Cl_2$) for the systematic covariate model and $Cl_0+Cl_1+Cl_2$ for the fixed allometric model, the elimination clearances of the metabolites (Cl_3 and Cl_4) and distribution volume of the central morphine compartment (V_1) versus bodyweight for the systematic covariate model (A) and the fixed allometric model (B).

Table IV. Bias (MPE) and precision (RMSE) of the predicted parameter values in the internal datasets stratified by age-group.

parameter	Age-group	Systematic covariate model		Fixed allometric model	
		MPE	RMSE	MPE	RMSE
Total morphine clearance	neonates [*]	4.26	0.426	10.8	0.922
	toddlers [†]	-0.74	1.65	12.7	5.00
	infants [‡]	4.63	3.28	-2.56	7.90
M3G clearance	neonates [*]	19.7	0.291	-19.5	2.50
	toddlers [†]	1.85	0.888	23.8	15.5
	infants [‡]	14.5	3.82	188	9.03
M6G clearance	neonates [*]	12.5	0.0784	-69.0	0.825
	toddlers [†]	0.262	0.230	100	5.27
	infants [‡]	4.93	0.852	253	2.84
Distribution volume of central morphine compartment	neonates [*]	23.2	1.30	90.6	2.93
	toddlers [†]	15.9	3.43	22.4	7.26
	infants [‡]	5.90	9.27	9.35	13.4

MPE = mean prediction error, RMSE = root mean square error,

^{*} neonates, PNA<30 days, n= 61,

[†] toddlers, PNA=1 month-1 year, n= 87,

[‡] infants, PNA>1 year n= 35

Table V. Bias (MPE) and precision (RMSE) of the predicted parameter values in the external datasets stratified by age-group.

parameter	age-group	systematic covariate model		fixed allometric model	
		MPE	RMSE	MPE	RMSE
Total morphine clearance	preterm neonates [*]	17.4	0.0667	192	0.444
	term neonates [†]	29.74	0.274	85.5	3.91
	toddlers [‡]	29.3	1.64	-26.6	4.22
M3G clearance	preterm neonates [*]	-16.3	0.102	114	0.225
	term neonates [†]	31.0	0.463	108	0.928
	toddlers [‡]	5.48	0.818	172	1.34
M6G clearance	preterm neonates [*]	-12.5	0.0447	46.8	0.779
	term neonates [†]	35.6	0.0818	31.1	0.502
	toddlers [‡]	12.5	0.285	93.5	0.490
Distribution volume of central morphine compartment	preterm neonates [*]	-12.9	0.945	-115	1.46
	term neonates [†]	-10.3	2.03	-296	3.50
	toddlers [‡]	3.45	3.47	-93.1	3.22

MPE = mean prediction error, RMSE = root mean square error

^{*} preterm neonates, (PNA < 30 days and PMA at birth < 36 weeks), n=80

[†] term neonates, (PNA < 30 days and PMA at birth ≥ 36 weeks), n=40

[‡] infants, PNA = 1 month – 1 year, n=33

8.4 Discussion

As in childhood many physiological changes take place in quick succession, the paediatric population is very heterogeneous. Additionally, studies in this population are often performed during routine clinical practice, which increases the variability in both dosing and sampling schemes, while due to limitations in sample size and frequency often only sparse data are obtained. All these factors influence the evaluation of population and covariate models for this young population. In the current study a framework of six different evaluation criteria is proposed for the evaluation of paediatric models. Most tools in the framework are not necessarily new, but in the context of paediatric model evaluation adaptations to the standard methods are sometimes required or a shift in emphasize on the various tools is essential.

As an example, two previously published paediatric population PK models for morphine that were based on the same dataset but fundamentally different covariate models, were evaluated with this framework. The systematic covariate model was developed by regarding bodyweight and age as regular covariates in a systematic covariate analysis (Chapter 3). The fixed allometric model was based on allometric principles including bodyweight *a priori* using exponential functions with fixed exponents and estimating an age-based function ^[3].

In itself the number of parameters in a model is not an evaluation criterion, however according to the rule of parsimony a model should have the lowest possible number of parameters. Large deviations of bootstrap parameter values from the original value, low precision in parameter estimates as expressed by high RSE values in the model fit and bootstrap procedures, small number of successful bootstrap runs and a condition number higher than 1000 are all generally indicative of model inaccuracy, model instability, and ill-conditioning ^[15-18]. Testing covariates for significance and only retaining a covariate when it significantly improves the model adheres to the rule of parsimony, does not introduce specific assumptions into the model, and the resulting model is always supported by data. In the current example, it is shown that the systematic covariate model performs well on all the criteria of ill-conditioning. In case covariates in a model are not formally tested for significance there is a risk of over-parameterization. The results for the fixed allometric model in this example suggest the fixed allometric model to be an unstable model for which precise parameter estimates cannot be obtained. It should also be noted in this respect that with the increased number of fixed and random model parameters in the fixed allometric model, the degrees of freedom in this model are increased as well. More degrees of freedom improve the description of data by a model, but do not necessarily improve the predictions by that model.

As stated before, paediatric PK analyses are often based on sparse data. This is an important aspect to consider when evaluating paediatric population PK models, because when data are uninformative due to the scarcity of the data, the estimated variability parameters may shrink to zero causing the individual *post hoc* parameter estimates to move towards the population predictions. This shrinkage phenomenon makes individual parameter estimates and the diagnostics based on them less reliable or even misleading ^[12,19]. Population predictions however, are solely based on the fixed effects described in the structural and covariate model. As the random effects are not considered in the population predictions, diagnostics based on population predictions are not sensitive to shrinkage and therefore more reliable than diagnostics based on individual predictions. The two models in the current study illustrate how diagnostics based on individual predictions can be misleading when shrinkage is high, as, according to table III, was the case for some of the parameters in both models. The plots of the individual predicted *versus* observed concentrations in figure 2 inadvertently suggest the fixed allometric model to perform better than the systematic covariate model, since especially for morphine the data points are closer to the line of unity. However, in figure 3 large differences in the predictive performance between the two models are revealed for both the internal and external datasets when considering population predicted concentrations instead of the individual predicted concentrations. For the systematic covariate model there are no signs of model misspecification. The fixed allometric model on the other hand shows significant bias towards under-prediction throughout the total concentration range. Since diagnostics based on population predictions are generally more reliable, these should always be included in the evaluation of paediatric (covariate) models.

Simulation-based diagnostics known as posterior predictive checks (PPC) are diagnostics that create a reference distribution of an observation of interest by performing multiple model simulations with inclusion of both fixed and random effects and subsequently compare the actual observations to this reference distribution ^[20]. A visual predictive check (VPC) is a commonly used and easily interpretable form of a PPC that graphically presents the reference distribution and observed data ^[21]. A VPC can be used for the evaluation of paediatric models as well, however when data are obtained during routine clinical practice and variability in individual dosing and sampling schemes are high, the NPDE methodology ^[13] is often easier to perform and interpret. Shrinkage does not influence the results of simulation-based diagnostics ^[12]. The results of the NPDE analysis of the models in the current example demonstrate that the systematic covariate model can quite accurately predict median concentrations for morphine and the glucuronides, but that it slightly over-predicts the variability in the overall dataset. This over-prediction of the variability is constant over time and over the concentration range. If this model

were to be used in simulation exercises the predictions would be unbiased and the inferences made on the variability in the population would be on the conservative side as the variability is predicted to be higher than it actually is. Based on the trend towards under-prediction by fixed allometric model, it can be concluded that significant bias in the predictions would occur if this model would be used in simulation exercises. The under-prediction of concentrations by the fixed allometric model increases the risk of overdosing when deriving morphine doses based on this model. These NPDE results substantiate the results obtained in the population predicted *versus* observed plots in figure 3.

Due to the heterogeneity in the paediatric population it is very important to not only perform diagnostics on the population as a whole, but to also look at various subpopulations by stratifying the datasets based on bodyweight or age.

For both models in the example, stratification showed the same descriptive and predictive performance in all age-groups. For the systematic covariate model the predictive performance of the model was adequate in all age-ranges. The trends towards under-prediction identified for the fixed allometric model was also similar across all age-ranges. Despite the fact that stratification of the diagnostics did not reveal new information in the current examples, this adjustment of the various validation tools remains imperative for the detection of previously unidentified age-related misspecifications.

To corroborate the obtained covariate relationships in paediatric models, the plots of individual and population parameter values *versus* the covariate presented here in figure 5, together with a numerical representation of bias and precision in table IV, have proven to be highly informative. Even in case of high shrinkage this diagnostic will enable the identification of bias in the population predictions of parameters. In this study, in both models bodyweight was the most important covariate for clearances and distribution volumes. For the systematic covariate model, population predicted parameter values are adequately centred in the range of individual predicted values for all parameters and across the entire bodyweight- and age-range. However, for the fixed allometric model, the population predicted parameter values are biased compared to the individual predicted values. For some parameters this bias exists over the total weight and age-range, for others only over part of these ranges. The results in the plots in figure 5 and table IV provide an explanation for the adequate individual concentration predictions by the fixed allometric model and the highly biased population predictions by this model, as observed with the other evaluation tools. Structural model misspecifications in the population parameter values result in biased population predictions that are corrected by the error models to yield good individual predictions. This type of diagnostic is hardly

ever published, however the information contained in these plots and tables is crucial and should become a standard diagnostic tool with paediatric population PK models.

Especially in drug development, population PK models in paediatric subpopulations are often used for extrapolations to younger age-ranges. When a model is used for this purpose, the obtained covariate relationships should be thoroughly evaluated, for instance by using the diagnostic in figure 5 and table IV. In the current study, two out of five external datasets (Ext 4 and Ext 5) include preterm patients, a younger and smaller population than the population in the internal dataset. For both models the results of the various tools in the framework were similar irrespective of the age-range in the dataset that was used, indicating that the inclusion of a new patient population in the external validation did not influence the overall results

A systematic covariate analysis is a data driven approach, therefore the extrapolation potential of the resulting model cannot be known *a priori*. It is clear that for the systematic covariate model in the current analysis extrapolations to older (heavier) children is not possible as the bodyweight based exponential equation for clearance predicts rapidly increasing clearances at higher weight-ranges. Figure 5 indicates that for this model, population parameter predictions are unbiased in the lower weight-ranges, suggesting that extrapolation to smaller children could be possible. The extensive evaluation procedures in the current example prove this to be the case in this particular example.

It is claimed that the allometric equations used in the fixed allometric model are based on 'sound biological principles' [22] and that the methodology based on these equations therefore yields mechanistic models that can be used for extrapolations outside the studied age- or weight-range. It is argued that the influence of size (parameterized by bodyweight) and maturation (parameterized by age) on the parameters in paediatric population PK models are disentangled by using the fixed allometric equations augmented by age-based functions [4,23]. However, the theory of allometry is based on the empirical observation that over a wide weight-range, metabolic rates of animal species increase with bodyweight to the power of 0.75 [24]. The fixed allometric exponents have no biological or physiological meaning, although reports exist that propose possible physiological explanations [25-27]. Conversely, a large body of evidence exists against the existence of one unique value for the allometric clearance exponent [28-34] and against the application of these allometric equations in human paediatric PK models [35,36]. Additionally, the maturation function based on age, only reflects a mathematical residue of the age-effect that remains after the inclusion of the correlated covariate bodyweight. In the current study, in the fixed allometric model the model-predicted increase in clearance comes to a plateau with increasing bodyweight. Therefore, it can not be excluded that

this model can be used for extrapolations to higher weight-ranges. The prospective properties of this model in preterm neonates is however very poor, albeit comparable to the predictions in the older children that comprised the learning dataset. As a result, the extrapolation potential of fixed allometric models in general can neither be confirmed nor disputed based on the results in the current study.

8.5 Conclusion

The framework of six evaluation criteria proposed in the current study takes into consideration the specific issues encountered in the evaluation of paediatric population models. The application of this framework to two models for morphine and its two major metabolites in children younger than three years with fundamentally different covariate models demonstrates how to detect over-parameterization, which is a risk with models based on sparse data. Additionally it illustrates the importance of diagnostics that are based on population predictions rather than individual predictions, as high shrinkage due to sparse data may yield misleading individual prediction-based diagnostics. Finally, the diagnostic comparing population parameter predictions with individually observed parameter values proved to be highly informative in assessing obtained covariate relationships as in the current example it detected the cause of model misspecification by the fixed allometric model. Stratification of the various diagnostics did not yield much additional information in the current examples, however due to the heterogeneity of the paediatric population this adaption of standard validation tools may be of value for other paediatric models.

The differences observed in model performance between the systematic covariate model and the fixed allometric model in the current study do not imply that any of the two methodologies for covariate model development is superior over the other. The current study does however highlight the importance of corroborating results in evaluation procedures. It also illustrates that information in data should not be ignored and that one should never be guided by theories alone.

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