

Personalized potassium and phosphorus intake recommendations can be reasoned for end-stage renal failure patients by simulating with hierarchical Bayesian multivariate model

Supplementary Materials

JARI TURKIA^{1,2} URSULA SCHWAB^{3,4} VILLE HAUTAMÄKI^{1,5}

¹ School of Computing, University of Eastern Finland, 80101 Joensuu, Finland

² CGI Suomi, Joensuu, Finland

³ School of Medicine, Institute of Public Health and Clinical Nutrition, University of Eastern Finland, Kuopio, Finland

⁴ Department of Medicine, Endocrinology and Clinical Nutrition, Kuopio University Hospital, Kuopio, Finland

⁵ Department of Electrical and Computer Engineering, National University of Singapore, Singapore

This notebook reproduces in detail the analysis of personal diet recommendations described in the main article. The notebook starts visualizing the collected raw data, prepares it for analysis and estimates personal reaction models with these data. The personal reaction models are combined with the estimation of the current personal diets for constructing personal graphical models. These personal models generate the levels of blood concentrations when a diet is given, they are used in simulating recommended personal diets for reaching predefined normal concentrations. Finally, these personal diet recommendations are compared for showing the divergence among the studied patients.

The notebook execution generates all the figures and tables that are included in the article, and produces also the referenced supplementary figures. The article is accompanied with a PDF rendition of the notebook that shows all the supplementary figures and important parts of the program code so that the analysis can be followed in detail. The executable RMarkdown notebook with data can be found in a public Github repository of the corresponding author.

Dialysis patient data

The analyzed dataset consists of food records and laboratory measurements from end-stage renal patients in dialysis. Following nutrients are possible considered as predictors of concentrations. Note that we use energy-% of fats and protein as unit in our analysis. This is referred as Table 1 in the main article.

Supplementary Table S 1: Nutrient predictors of the model.

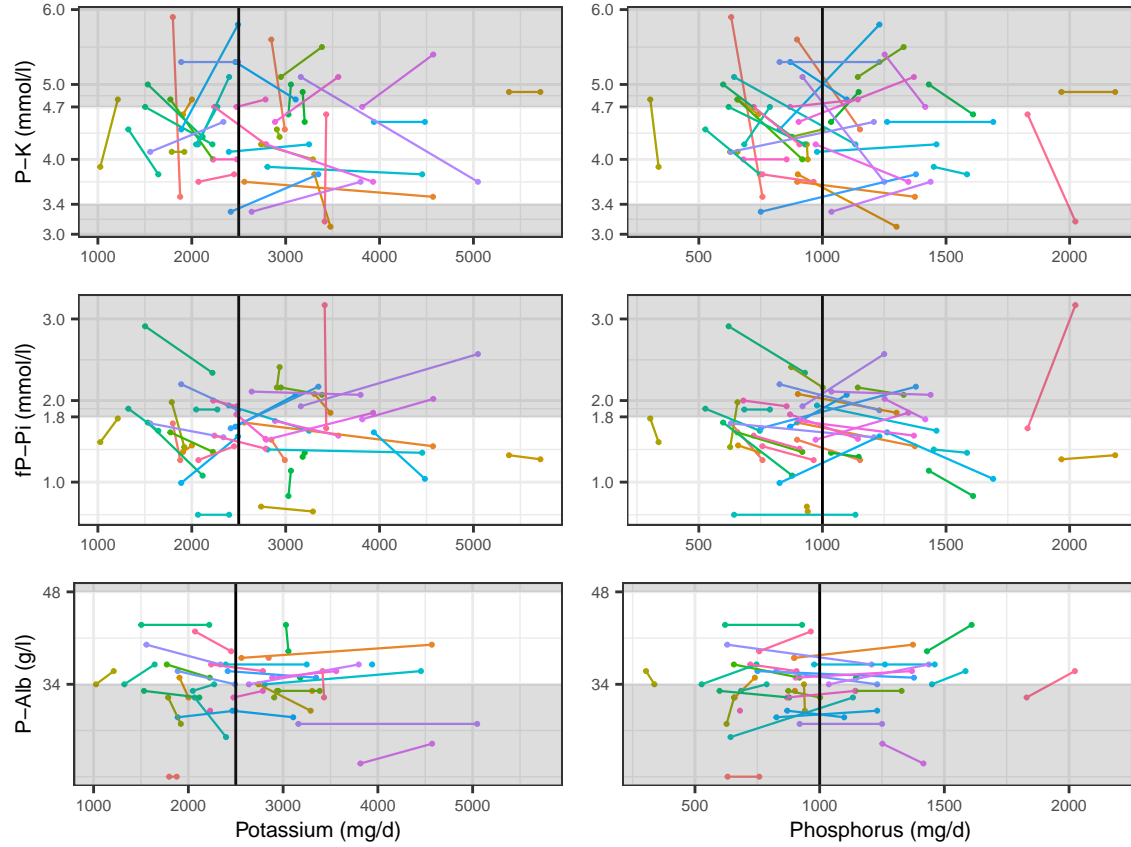
Nutrient	Sample avg. (min-max)
Carbonhydrates, E%	43.6 (27.1 - 63.6) E%
Fat E%	38.9 (23.4 - 54.1) E%
Monounsaturated Fatty Acids, E%	14.7 (5.6 - 25.1) E%
Polyunsaturated Fatty Acids, E%	7.1 (2.2 - 15.8) E%
Protein, E%	15.1 (9.2 - 22.4) E%
Saturated Fatty Acids, E%	13.7 (5.9 - 24.5) E%
Fiber	17 (5 - 42) g/d
Protein, g/kg	0.8 (0.2 - 2.1) g/kg/d
Energy, kcal/kg	21.8 (5.6 - 58.6) kcal/kg
Calsium	570 (123 - 1741) mg/d
Natrium	2588 (813 - 5487) mg/d
Phosphorous	1042 (304 - 2184) mg/d
Potassium	2785 (1026 - 5713) mg/d
Salt	6560 (201 - 13863) mg/d
Water	1804 (601 - 3613) ml/d
Vitamin D	8 (0 - 31) ug/d

and also following personal details and medication are considered as predictors. This is referenced as Table 2 in the article.

Supplementary Table S 2: Personal details that are used as predictors.

Personal detail	Percentage of patients
Blood fat medication	68%
Diabetes medication	51%
Hydroxycholecalciferol	43%
Phosphate binder med.	22%
Renavit	97%
Gender	41% female
Home hemodialysis	24%
Hospital hemodialysis	57%
Peritoneal dialysis	19%

Let us explore then how the intake of dietary potassium and phosphorus corresponds in data to plasma concentrations of potassium, phosphate and albumin



Supplementary Figure S 1: Figure shows the progress between two observations of plasma potassium (P-K), fasting plasma phosphate (fP-Pi) and plasma albumin (P-Alb) concentrations of the studied patients. White regions in the panels show the recommended concentration levels, P-K 3.4 - 4.7 mmol/l, fP-Pi < 1.8 mmol/l and P-Alb 34 - 48 g/l. Vertical black lines denote the commonly recommended maximum intakes of these nutrients. The goal is to find personal intake levels that keep the concentrations in recommended levels, if possible. The figure is plotted with ggplot2 package for R language (v 3.3.5, <https://ggplot2.tidyverse.org>).

Aim of this analysis is to find such personal levels of potassium and phosphorous intakes that keep all these concentrations in their normal levels marked with white regions, if possible.

Development of nutrition reaction model

In this work we construct personal generative models for concentrations that allow conditioning personal diet recommendations. Essential part of these models is the personal reactions to nutrients and other predictors. Simultaneous reactions on all considered concentrations are modeled as multivariate model that has all the concentrations are response variables.

For comparison, we estimate multivariate systems with and without cross-model covariance. With cross-model covariance estimated, the model corresponds to seemingly unrelated model system, and without the model is Bayesian network with separated local distributions. This simpler model is estimated first with only potassium and phosphorous concentrations as responses.

```
initial_graph <- mebn.fully_connected_bipartite_graph(datadesc_fat_epros)
```

```
pk_fppi_targets <- datadesc_fat_epros[datadesc_fat_epros$Name %in% c('pk','fppi'),]

dialdiet_gamma <- mebn.bipartite_model(reaction_graph = initial_graph,
                                     inputdata = dialysis,
                                     predictor_columns = assumedpredictors_fat_epros,
                                     assumed_targets = pk_fppi_targets,
                                     group_column = "potilas",
                                     local_estimation = mebn.sampling,
                                     local_model_cache = "models/BLMM_gamma_separate",
                                     stan_model_file = "mebn/v2/BLMM_gamma.stan",
                                     normalize_values = TRUE)

# Write the generated graph in a GraphML file
write.graph(dialdiet_gamma, "graphs/dialysis_gamma_separate.graphml", "graphml")
```

In Bayesian network both responses were estimated separately. Next, we estimate a multivariate model where both distribution are estimated during single sampling. It does not factorize into separate distributions, but allows using more data.

```
dialdiet_gamma_mv2_epros <- mebn.bipartite_multivariate(reaction_graph = initial_graph,
                                                       inputdata = dialysis,
                                                       predictor_columns = assumedpredictors_fat_epros,
                                                       assumed_targets = pk_fppi_targets,
                                                       group_column = "potilas",
                                                       local_estimation = mebn.multivariate_sampling,
                                                       local_model_cache =
                                                         "models/BLMM_gamma_qr_multivariate2/fat_epros",
                                                       stan_model_file = "mebn/v2/BLMM_gamma_qr_mv.stan",
                                                       normalize_values = TRUE)

write.graph(dialdiet_gamma_mv2_epros,
            "graphs/dialysis_gamma_multivariate2_epros.graphml", "graphml")
```

Next we like to add plasma albumin concentration as a third constraint in the model, but unfortunately 8 of 37 patients have missing albumin measurements. From such a small dataset we don't want to remove any patients and so we predict the missing albumin levels and impute them to dataset.

For prediction, estimate the model without missing values in P-Alb. The rows with missing values are held out from the density estimation, but patients are kept in the model for estimating their parameters.

```
pk_fppi_palb_targets <-
  datadesc_fat_epros[datadesc_fat_epros$Name %in% c('pk','fppi','palb'),]

# 0/1-index for palb = NA
holdout_index <- as.vector(as.numeric(is.na(dialysis$palb)))

# Stan does not support NA in data (in Y), so let's change NA to magic number
if (anyNA(dialysis$palb)) dialysis[is.na(dialysis$palb),]$palb <- -1

initial_graph <- mebn.fully_connected_bipartite_graph(datadesc_fat_epros)

dialdiet_gamma_mv3_missing_palb <- mebn.bipartite_multivariate(
  reaction_graph = initial_graph,
```

```

inputdata = dialysis,
targetdata = holdout_index,
predictor_columns = assumedpredictors_fat_epros,
assumed_targets = pk_fppi_palb_targets,
group_column = "potilas",
local_estimation = mebn.multivariate_sampling,
local_model_cache =
  "models/BLMM_gamma_qr_multivariate3/fat_epros_missing_palb",
stan_model_file = "mebn/v2/BLMM_gamma_qr_mv_cv.stan",
normalize_values = TRUE)

```

Personal data imputation models

Now we can create personal models for patients that have missing plasma albumin measurements. These models are then used for making personal predictions for replacing missing values.

```

# Extract personal generative models for patients who have missing P-Alb values

patients_with_missing_palb <- unique(as.vector(dialysis[is.na(dialysis$palb),]$potilas))

# - initial graph structure
initial_graph <- mebn.fully_connected_bipartite_graph(datadesc_fat_epros)

# - previously estimated graphical model with all the persons
local_distributions <- pk_fppi_palb_targets
local_distributions$modelcache <-
  "models/BLMM_gamma_qr_multivariate3/fat_epros_missing_palb"

# - get personal data, normalized and original

# - statistics for vertex levels are calculated from normalized data
predictors <- nrow(assumedpredictors_fat_epros)
normalized_input <- sapply(1:predictors, mebn.scale_gaussians,
  data = dialysis, datadesc = assumedpredictors_fat_epros)
normalized_input_df <- as.data.frame(normalized_input)

datadesc <- datadesc_fat_epros

```

Then we predict missing values with these data imputing models

```

for (i in 1:nrow(rows_with_missing_palb)) {

  datarow <- rows_with_missing_palb[i,]

  # get model for patient in this datarow

  personal_model_dir <- paste0("data_imputation_models/", datarow$potilas)
  print(paste0("Reading personal data imputation model ", personal_model_dir,
    "/personal_graph.graphml"))

  personal_graph <- read.graph(paste0(personal_model_dir, "/personal_graph.graphml"), "graphml")
}

```

```

# and use the data in this row to predict palb
evidence <- rows_with_missing_palb[i, assumedpredictors_fat_epros$Name]

posterior_prediction <- mebn.personal_prediction(reaction_graph = personal_graph,
  graph_dir = personal_model_dir,
  evidence = evidence,
  stan_model_file = "diet/posterior_prediction.stan")

# personal_predictions contains predictions multivariate predictions
# P-K and fP-Pi are known, but P-Alb is missing

posterior <- rstan::extract(posterior_prediction, par= "posterior[3]")

# use predicted posterior mean for missing P-Alb value
# - this is i:th NA value in dialysis
predicted_palb <- mean(posterior$posterior[3])

# rows in 'rows_with_missing_palb' and 'original_palb' are in same order
# so we can imput in the NA values at 'original_palb'

for (m in 1:imput_length)
{
  if (is.na(original_palb[m]))
  {
    original_palb[m] <- predicted_palb
    break;
  }
}
}

dialysis2 <- dialysis
dialysis2$palb <- original_palb
saveRDS(dialysis2, "data/DIALYSIS_imputed_palb.rds")

```

With this data imputation we can estimate a cross-covariance model with three responses

```

pk_fppi_palb_targets <- datadesc_fat_epros[datadesc_fat_epros$Name %in% c('pk','fppi','palb'),]

initial_graph <- mebn.fully_connected_bipartite_graph(datadesc_fat_epros)

dialysis_imputed <- readRDS("data/DIALYSIS_imputed_palb.rds")
no_holdout <- rep(0, nrow(dialysis_imputed))

dialdiet_gamma_mv3 <- mebn.bipartite_multivariate(reaction_graph = initial_graph,
  inputdata = dialysis_imputed,
  targetdata = no_holdout,
  predictor_columns = assumedpredictors_fat_epros,
  assumed_targets = pk_fppi_palb_targets,
  group_column = "potilas",
  local_estimation = mebn.multivariate_sampling,
  local_model_cache =
    "models/BLMM_gamma_qr_multivariate3/imputed_palb",

```

```

stan_model_file = "mebn/v2/BLMM_gamma_qr_mv_cv.stan",
normalize_values = TRUE)

write.graph(dialdiet_gamma_mv3,
            "graphs/dialysis_gamma_multivariate3_imputed.graphml", "graphml")

```

Finally, we consider also the effect of dialysis treatment type as new level of grouping in data. This allows us to estimate average effects of nutrition for each treatment type and then also personal effects within those treatments.

```

dialysis_imputed <- readRDS("data/DIALYSIS_imputed_palb.rds")

# add the dialysis treatment type as a grouping factor
dialysis_imputed$hoitoryhma <- as.factor(dialysis_imputed$hoitomuoto)

# and sort the data by treatment/patient/observation
dialysis_imputed <- dialysis_imputed[order(dialysis_imputed$hoitoryhma,
                                           dialysis_imputed$potilas, dialysis_imputed$havainto),]

dialdiet_gamma_mv3_two_level <- mebn.bipartite_two_level_multivariate(
  reaction_graph = initial_graph,
  inputdata = dialysis_imputed,
  targetdata = no_holdout,
  predictor_columns = assumedpredictors_fat_epros,
  assumed_targets = pk_fppi_palb_targets,
  group_column = "hoitoryhma",
  subject_column = "potilas",
  local_estimation = mebn.two_level_multivariate_sampling,
  local_model_cache =
    "models/BLMM_gamma_mv_cross/two_levels",
  stan_model_file =
    "mebn/v2/BLMM_gamma_two_level_grouping.stan",
  normalize_values = TRUE)

write.graph(dialdiet_gamma_mv3_two_level,
            "graphs/dialysis_gamma_two_level_grouping.graphml", "graphml")

```

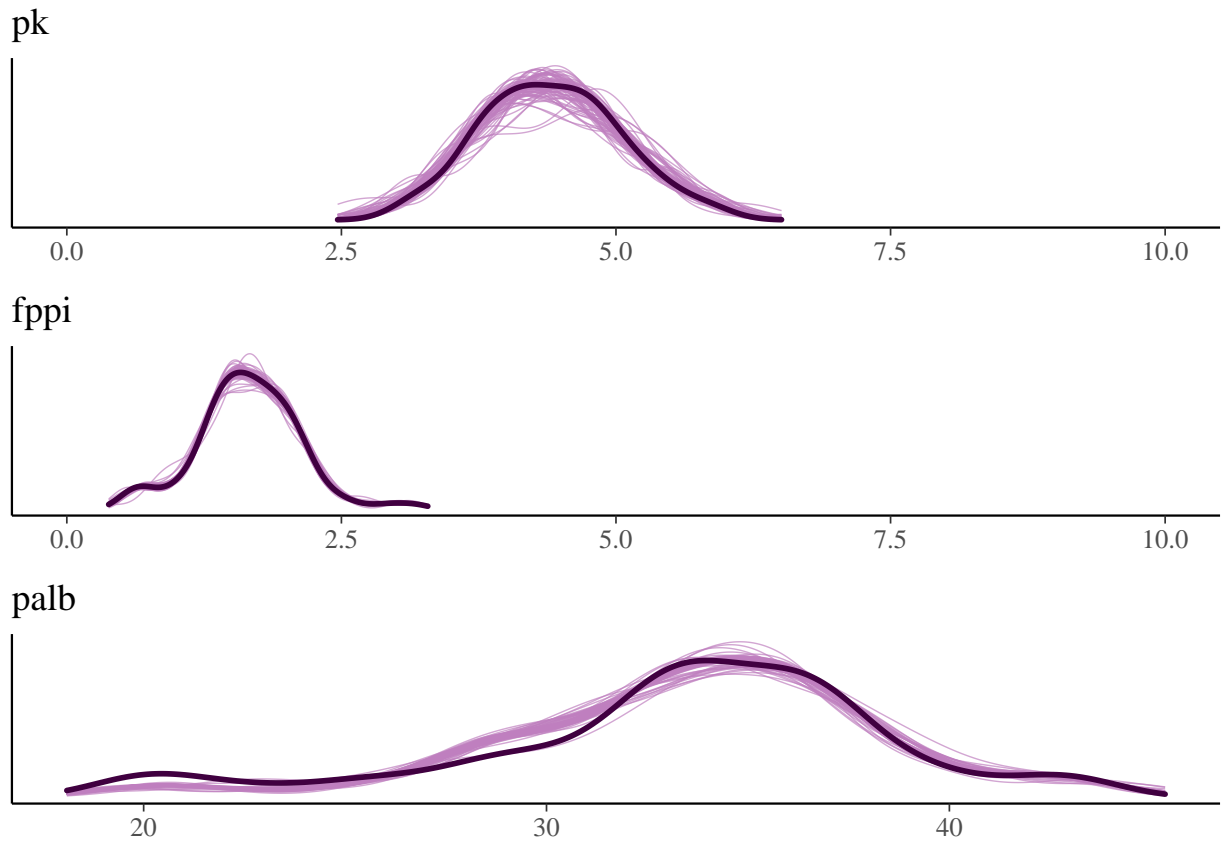
Model analysis

Here we evaluate the model's ability to repeat the measured observations with normalized root mean square error (NRMSE) and compare how adding the dialysis treatment as an explicit hierarchical layer in the model affects this error. We also asses the model's performance with visual posterior predictive check by comparing samples from the model with measured values. This would give away possible bias in the model.

Supplementary Table S 3: Personal graphical models with different reaction model candidates are evaluated with NRMSE for each modelled concentration and average model error.

Reaction model	NRMSE			
	P-K	fP-Pi	P-Alb	average error
mv3_cross_two_levels	0.004	0.007	0.002	0.004
mv3_cross_single_level	0.036	0.13	0.059	0.036

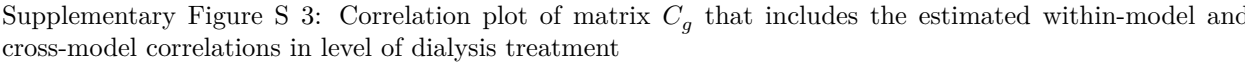
```
## [1] "pk"
## [1] "fppi"
## [1] "palb"
```

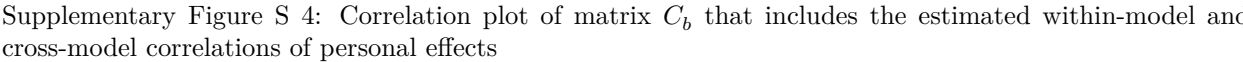


Supplementary Figure S 2: Posterior predictive check of the model where three concentrations and their parameters are stacked into one univariate model for estimating cross-model correlations.

Then we drill into the estimated correlation matrices of effects. The model estimates how much the effects of nutrients vary from general effects in dialysis treatment level and further how the effects vary personally within a treatment. These correlation plots show how the effects are correlated.

```
## Scale for 'fill' is already present. Adding another scale for 'fill', which
## will replace the existing scale.
```



Following tables include all the estimated nutrition effects in general, dialysis treatment and personal levels

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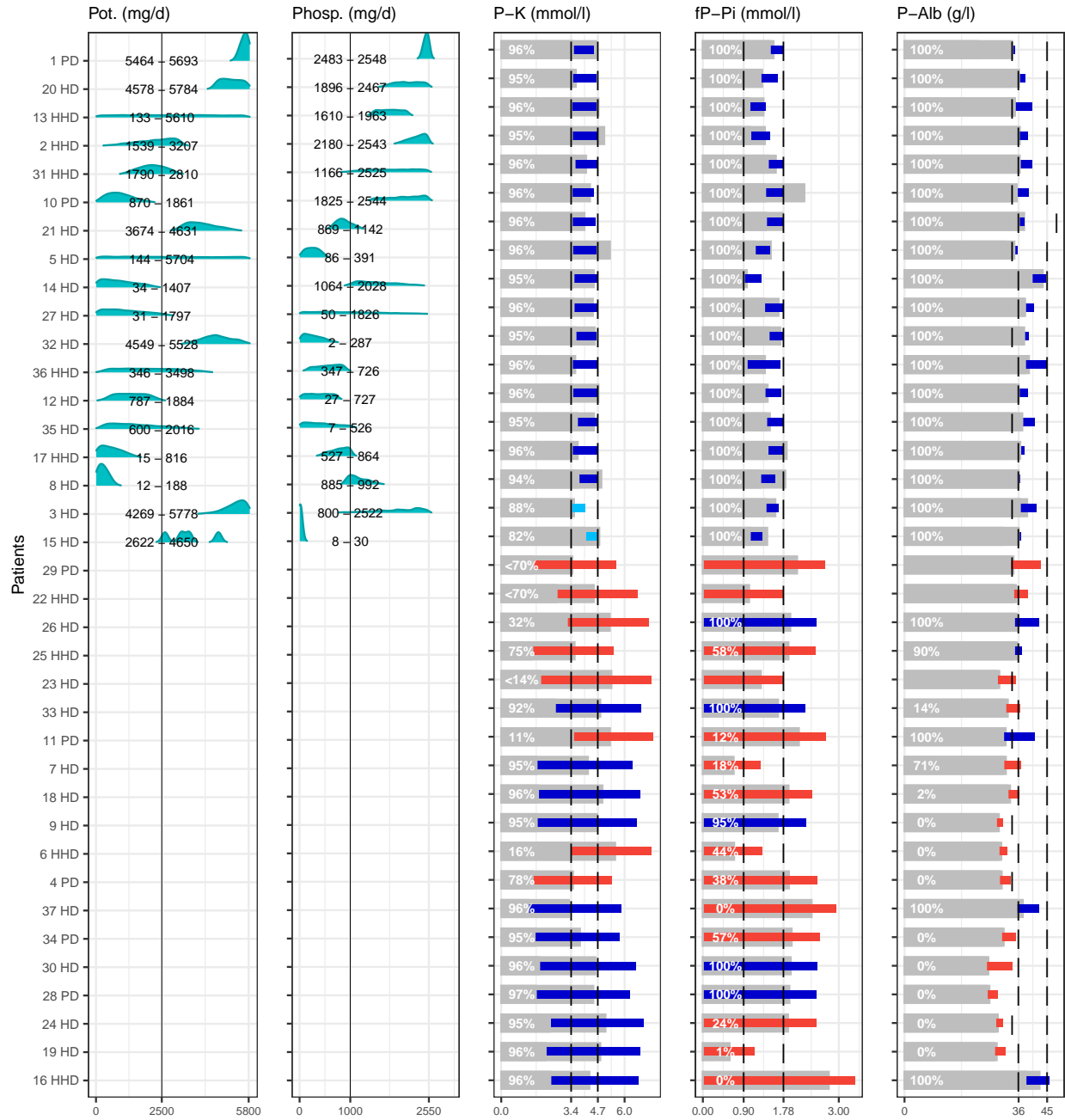
personal effects are shown within each treatment ($\hat{\beta}_{ij} + \hat{g}_{ijk} + \hat{b}_{ijp}$). The table is sorted in decreasing order of between-treatment variation ($\hat{\sigma}_g$) and all the estimates include their 90%-credible intervals.

Nutrient	Conc.	General effect	Home hemodialysis			Hospital hemodialysis			Peritoneal dialysis		
			avg	min	max	avg	min	max	avg	min	max
Water	P-Alb	-0.94 [-1.03; 2.79]	-1.33 [-3.96; 2.19]	-2.09 [-5.21; 2.75]	-0.99 [-4.14; 2.37]	-0.43 [-2.51; 1.46]	-0.77 [-2.17; 1.47]	0.36 [-3.68; 3.52]	-1.53 [-5.03; 3.24]	-2.08 [-5.59; 3.46]	-0.97 [-5.15; 3.14]
Natrium	P-Alb	-2.60 [-0.24; 7.58]	-0.92 [-9.95; 7.96]	-2.78 [-16.47; 7.74]	0.21 [-6.29; 8.56]	-0.27 [-7.82; 7.72]	-1.87 [-13.40; 7.77]	1.34 [-4.12; 8.06]	-6.70 [-30.75; 7.59]	-7.97 [-36.19; 8.14]	-5.36 [-24.93; 7.70]
Energy, kcal/kg	P-Alb	3.46 [1.76; 15.59]	2.63 [-5.85; 12.96]	1.07 [-6.48; 11.89]	3.59 [-5.90; 14.48]	3.42 [-5.54; 14.49]	1.76 [-7.86; 12.96]	4.16 [-4.11; 15.21]	3.49 [-5.50; 15.16]	2.91 [-5.71; 15.90]	4.13 [-4.85; 17.20]
Phosphorous	P-Alb	0.24 [0.30; 9.18]	0.83 [-6.93; 7.38]	-0.40 [-8.26; 7.29]	1.45 [-6.89; 8.04]	-0.87 [-5.55; 6.58]	-2.26 [-8.24; 6.32]	0.61 [-5.61; 7.50]	0.85 [-5.63; 9.97]	-0.06 [-6.80; 9.46]	1.79 [-5.14; 11.47]
Diabetes medication	P-Alb	0.98 [2.55; 5.86]	-2.22 [-8.21; 7.17]	-2.99 [-9.13; 6.95]	-1.75 [-9.01; 8.29]	0.99 [-5.89; 4.66]	0.01 [-7.55; 4.72]	1.73 [-6.02; 6.63]	0.99 [-7.53; 10.12]	-0.28 [-10.08; 9.25]	1.26 [-8.79; 10.17]
Renavit	P-Alb	7.67 [10.56; 23.52]	8.89 [-11.49; 22.52]	8.78 [-12.02; 22.63]	9.72 [-11.07; 23.87]	5.50 [-10.54; 22.08]	4.28 [-10.88; 22.95]	6.86 [-10.28; 22.56]	6.41 [-11.66; 22.37]	6.07 [-12.09; 22.46]	7.41 [-11.94; 23.58]
Hydroxycholecalciferol	P-Alb	3.73 [4.17; 10.25]	2.74 [-7.13; 9.43]	0.88 [-9.11; 9.33]	2.86 [-9.90; 11.57]	3.01 [-4.00; 9.51]	1.68 [-6.12; 7.45]	4.57 [-4.02; 12.35]	2.26 [-9.21; 9.34]	1.19 [-10.08; 9.84]	3.46 [-11.16; 14.06]
Potassium	P-Alb	-0.33 [-0.22; 3.86]	-0.97 [-6.09; 3.72]	-2.20 [-6.85; 3.35]	0.45 [-7.04; 7.24]	0.22 [-2.90; 3.23]	-1.41 [-4.22; 2.72]	1.48 [-1.88; 5.78]	0.90 [-3.16; 5.94]	0.12 [-4.47; 5.37]	1.85 [-4.13; 7.86]
Phosphate binder med.	P-Alb	-1.17 [-2.86; 14.60]	-0.19 [-9.29; 10.80]	-2.12 [-9.90; 9.54]	1.38 [-10.25; 9.85]	-1.18 [-9.01; 6.76]	-4.28 [-12.97; 7.67]	0.51 [-9.86; 8.53]	-0.40 [-7.38; 8.54]	0.58 [-9.27; 10.76]	1.29 [-7.68; 10.11]
Carbonhydrates, E%	P-Alb	-0.08 [-1.17; 10.36]	0.12 [-12.32; 11.02]	0.01 [-13.04; 11.41]	0.97 [-12.77; 11.32]	-0.42 [-13.22; 9.82]	-1.34 [-13.94; 9.83]	0.67 [-13.89; 10.27]	-0.39 [-13.16; 10.13]	-1.57 [-14.61; 10.58]	-0.38 [-12.88; 9.79]
Fiber	P-Alb	0.67 [1.37; 4.02]	0.74 [-3.17; 3.45]	0.62 [-3.62; 3.69]	0.96 [-3.23; 4.63]	0.89 [-2.88; 3.37]	0.63 [-3.18; 3.47]	1.25 [-3.14; 4.96]	0.54 [-5.02; 5.33]	0.49 [-4.98; 5.52]	0.66 [-5.10; 5.86]
Gender	P-Alb	-1.11 [0.04; 7.33]	-0.82 [-13.78; 6.85]	-2.29 [-17.44; 6.89]	0.30 [-15.40; 8.63]	2.62 [-5.38; 9.55]	1.14 [-7.53; 9.67]	4.31 [-4.49; 16.23]	-1.74 [-13.81; 9.41]	-3.39 [-15.65; 7.52]	-0.90 [-13.88; 14.29]
Blood fat medication	P-Alb	-1.17 [-0.48; 9.57]	0.65 [-6.05; 13.04]	0.06 [-7.70; 13.61]	1.68 [-6.12; 15.32]	-1.66 [-5.63; 4.27]	-2.85 [-9.07; 4.49]	-0.36 [-5.07; 5.04]	1.96 [-6.75; 11.86]	1.50 [-8.54; 11.88]	2.29 [-7.17; 11.72]
Blood fat medication	P-K	-1.42 [-1.74; 2.05]	-0.22 [-3.30; 3.06]	-0.30 [-3.64; 3.11]	0.02 [-2.97; 3.25]	-0.41 [-1.33; 0.90]	-0.53 [-1.69; 0.94]	-0.25 [-1.41; 1.35]	-0.42 [-4.45; 1.92]	-0.58 [-5.03; 1.82]	-0.27 [-4.20; 1.92]
Diabetes medication	P-K	-0.77 [-0.09; 1.45]	-1.04 [-3.70; 0.93]	-1.17 [-4.67; 1.07]	-0.62 [-3.47; 1.17]	-0.40 [-1.97; 0.73]	-0.75 [-3.32; 0.61]	-0.16 [-1.99; 1.31]	-1.64 [-3.86; 1.49]	-1.85 [-4.07; 1.40]	-1.39 [-3.59; 1.52]
Gender	P-K	-0.37 [-0.33; 2.45]	0.35 [-1.18; 2.54]	-0.32 [-3.71; 2.80]	0.81 [-1.06; 3.10]	0.20 [-1.35; 1.39]	-0.35 [-1.89; 1.65]	0.60 [-1.34; 2.32]	0.97 [-1.19; 2.77]	0.79 [-1.51; 3.33]	1.34 [-1.38; 3.52]
Carbonhydrates, E%	P-Pi	0.27 [0.21; 3.28]	-0.07 [-1.87; 2.32]	-0.10 [-1.87; 2.34]	-0.05 [-1.82; 2.34]	0.42 [-1.05; 2.66]	0.38 [-1.13; 2.65]	0.45 [-1.04; 2.69]	0.19 [-2.02; 3.59]	0.10 [-2.10; 3.53]	0.16 [-1.98; 3.62]
Fat E%	P-K	-0.68 [-0.26; 3.60]	-0.71 [-4.74; 3.07]	-0.85 [-6.11; 3.03]	-0.49 [-3.53; 3.29]	-0.67 [-4.62; 4.02]	-0.95 [-5.00; 3.65]	-0.47 [-4.42; 4.62]	-0.68 [-5.02; 2.63]	-0.96 [-5.88; 2.50]	-0.58 [-5.22; 2.87]
Water	P-K	0.03 [0.23; 2.11]	-0.04 [-2.09; 0.97]	-0.12 [-2.39; 1.00]	0.07 [-2.23; 1.14]	0.12 [-0.44; 0.71]	0.07 [-0.53; 0.63]	0.20 [-0.58; 0.96]	-0.16 [-2.92; 1.43]	-0.24 [-3.02; 1.37]	-0.08 [-2.75; 1.61]
Renavit	P-K	2.46 [2.45; 6.63]	2.52 [-4.12; 8.39]	2.46 [-4.28; 8.64]	2.77 [-4.03; 9.24]	2.74 [-2.38; 6.89]	2.52 [-2.23; 6.82]	3.06 [-2.24; 7.19]	3.20 [-2.46; 8.22]	2.88 [-3.27; 7.77]	3.39 [-2.35; 8.18]
Protein, g/kg	P-Alb	-3.36 [-1.96; 4.65]	-6.42 [-15.03; 4.12]	-7.96 [-15.10; 4.31]	-4.16 [-13.91; 5.61]	-3.61 [-13.13; 4.26]	-6.61 [-13.09; 3.76]	-1.58 [-12.52; 6.91]	-2.80 [-17.04; 5.18]	-4.29 [-17.36; 5.47]	-1.43 [-18.18; 10.29]
Monounsaturated Fatty Acids, E%	P-Alb	-0.21 [-1.41; 7.64]	-0.81 [-5.32; 6.33]	-1.56 [-7.19; 6.64]	0.30 [-5.38; 6.36]	0.65 [-5.21; 6.08]	-1.04 [-6.26; 6.10]	2.12 [-5.80; 6.43]	-0.32 [-5.46; 6.88]	-1.23 [-6.28; 7.11]	0.97 [-5.46; 7.37]
Calcium	P-Alb	1.71 [2.37; 7.24]	1.20 [-2.27; 6.36]	0.47 [-4.00; 5.89]	1.62 [-2.66; 8.20]	-0.01 [-4.83; 3.66]	-0.41 [-5.41; 3.16]	0.71 [-3.47; 4.53]	0.96 [-4.96; 6.27]	0.57 [-6.78; 6.68]	1.18 [-5.96; 7.21]
Fat E%	P-Alb	1.30 [3.41; 12.51]	1.82 [-11.70; 12.55]	1.02 [-12.73; 12.42]	2.89 [-11.07; 13.62]	1.77 [-11.60; 12.42]	1.02 [-12.90; 12.69]	3.21 [-10.96; 12.70]	2.29 [-11.90; 13.25]	2.10 [-12.15; 13.13]	3.09 [-10.81; 13.83]
Vitamin D	P-Alb	1.02 [1.80; 4.83]	1.38 [-1.70; 3.73]	0.97 [-3.84; 4.93]	2.03 [-1.37; 5.05]	1.26 [-0.58; 2.48]	0.99 [-1.99; 3.09]	1.66 [-0.63; 4.42]	1.19 [-2.02; 5.15]	0.89 [-3.11; 5.20]	1.40 [-2.86; 6.39]
Phosphorous	P-K	0.18 [0.06; 1.91]	-0.16 [-1.65; 1.54]	-0.37 [-2.92; 1.57]	0.08 [-1.29; 2.18]	0.24 [-1.20; 1.60]	0.02 [-2.11; 1.51]	0.60 [-0.57; 2.58]	-0.15 [-2.42; 1.70]	-0.51 [-3.66; 1.78]	0.09 [-2.12; 2.07]
Hydroxycholecalciferol	P-K	-2.20 [-0.70; 3.33]	-1.44 [-5.94; 3.29]	-1.63 [-6.11; 3.32]	-1.23 [-6.09; 4.08]	-0.23 [-1.55; 1.43]	-0.56 [-2.90; 1.45]	0.27 [-1.65; 3.95]	-1.41 [-5.15; 3.91]	-1.79 [-5.03; 3.53]	-1.29 [-4.99; 4.80]
Polysaturated Fatty Acids, E%	P-Alb	-2.95 [-3.20; 2.81]	-1.50 [-5.03; 4.81]	-1.87 [-5.58; 4.18]	-1.12 [-5.05; 6.15]	-3.06 [-5.82; 0.19]	-3.49 [-5.85; -0.32]	-2.55 [-5.83; 2.79]	-1.91 [-4.94; 2.78]	-2.16 [-6.19; 3.18]	-1.61 [-4.83; 3.58]
Salt	P-Alb	7.19 [-1.91; 37.18]	1.87 [-8.54; 14.87]	1.48 [-9.11; 14.84]	2.14 [-8.25; 14.92]	1.30 [-5.87; 7.79]	0.76 [-9.44; 7.59]	1.99 [-3.85; 7.92]	3.87 [-9.82; 27.21]	3.68 [-10.66; 27.11]	4.37 [-9.20; 27.28]
Energy, kcal/kg	P-K	-0.23 [-1.01; 5.76]	1.00 [-2.34; 5.98]	0.80 [-2.42; 4.79]	1.17 [-2.42; 5.21]	-0.53 [-2.72; 4.53]	-0.89 [-3.12; 2.80]	-0.35 [-2.86; 6.97]	-0.75 [-4.05; 5.44]	-0.96 [-4.03; 4.68]	-0.56 [-3.88; 6.43]
Natrium	P-K	-1.94 [-0.62; 1.53]	-1.36 [-8.97; 1.74]	-1.67 [-9.14; 1.61]	-0.90 [-8.30; 1.95]	-0.57 [-1.57; 0.71]	-0.96 [-1.88; 0.66]	-0.23 [-1.57; 0.96]	-1.74 [-5.59; 1.06]	-2.18 [-6.16; 1.14]	-1.68 [-5.51; 1.18]
Saturated Fatty Acids, E%	P-Alb	-3.58 [-5.11; 5.14]	-2.69 [-7.00; 5.14]	-2.91 [-7.55; 4.83]	-2.38 [-6.77; 5.28]	-2.15 [-6.96; 2.74]	-2.72 [-8.62; 2.46]	-1.74 [-6.70; 3.49]	-1.81 [-7.80; 9.35]	-1.88 [-8.43; 9.87]	-1.35 [-7.35; 9.49]
Hydroxycholecalciferol	P-Pi	0.27 [0.27; 2.94]	-0.16 [-1.67; 2.17]	-0.33 [-2.38; 2.31]	0.09 [-1.58; 3.70]	0.50 [-0.58; 1.66]	0.13 [-1.72; 1.81]	0.77 [-0.44; 1.84]	0.10 [-2.28; 1.77]	-0.17 [-3.07; 2.03]	0.28 [-2.39; 2.21]

Supplementary Table S4: Nutrition effect magnitudes from nutrients and other modeled features ($j = 1, \dots, 22$) to blood concentrations ($i = 1, \dots, 3$) for analyzed patients ($p = 1, \dots, 37$) in all three additive levels of the model. General effects ($\hat{\beta}_{ij}$) are shown to vary between patients in home hemodialysis, hospital hemodialysis, and peritoneal dialysis. First column of each dialysis type (avg) shows typical effect of the treatment ($\hat{\beta}_{ij} + \hat{g}_{ijk}$, $k = 1, \dots, 3$) that can further vary personally. Minimum and maximum of these personal effects are shown within each treatment ($\hat{\beta}_{ij} + \hat{g}_{ijk} + \hat{b}_{ijp}$). The table is sorted in decreasing order of between-treatment variation ($\hat{\sigma}_g$) and all the estimates include their 90%-credible intervals (table continues).

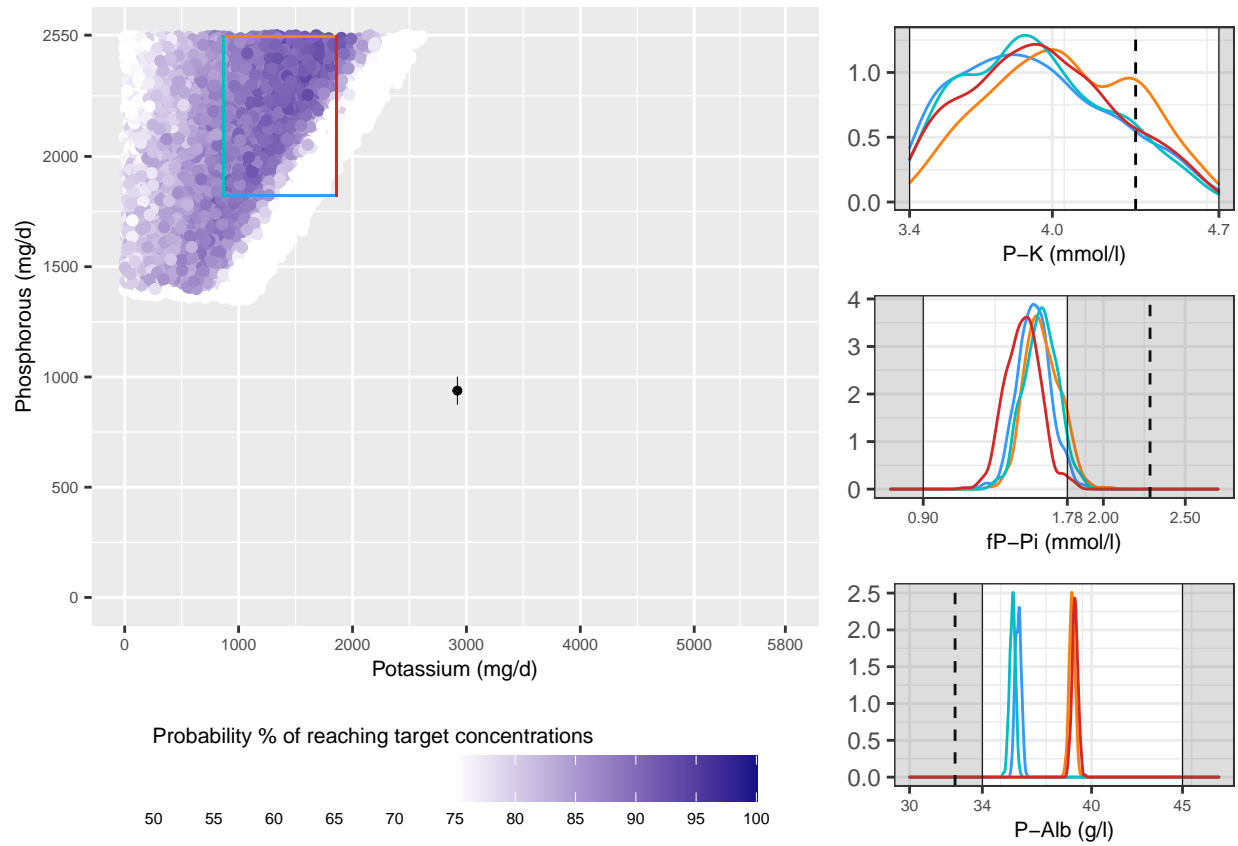
Nutrient	Conc.	General effect	Home hemodialysis			Hospital hemodialysis			Peritoneal dialysis		
			avg	min	max	avg	min	max	avg	min	max
Protein, E%	P-Alb	2.61 [2.41; 9.62]	1.43 [-4.90; 5.88]	1.05 [-5.71; 6.62]	2.68 [-4.73; 7.60]	1.85 [-4.02; 6.45]	0.95 [-5.27; 5.72]	2.76 [-3.46; 8.24]	2.57 [-5.36; 10.19]	1.79 [-6.12; 8.54]	2.88 [-4.90; 10.60]
Fiber	fl-Pi	0.49 [0.10; 2.28]	0.12 [-0.61; 0.82]	0.04 [-0.75; 0.61]	0.16 [-0.66; 0.78]	-0.15 [-0.69; 0.51]	-0.26 [-0.86; 0.49]	-0.09 [-0.60; 0.55]	-0.11 [-0.73; 0.99]	-0.16 [-0.80; 0.85]	-0.07 [-0.80; 0.99]
Polyunsaturated Fatty Acids, E%	P-K	-0.05 [0.00; 1.43]	-0.25 [-1.47; 0.73]	-0.35 [-1.50; 0.77]	-0.14 [-1.42; 1.10]	-0.09 [-0.93; 0.53]	-0.20 [-1.19; 0.57]	-0.01 [-0.97; 0.90]	0.02 [-0.97; 1.06]	-0.05 [-0.98; 0.83]	0.07 [-1.07; 1.09]
Monounsaturated Fatty Acids, E%	P-K	1.62 [1.27; 4.01]	1.26 [-1.07; 3.33]	1.06 [-1.22; 2.63]	1.55 [-0.94; 3.59]	0.67 [-0.81; 1.85]	0.28 [-1.62; 1.65]	0.92 [-0.92; 2.24]	0.38 [-1.28; 2.10]	-0.22 [-2.35; 1.53]	0.47 [-1.06; 2.53]
Carbonhydrates, E%	P-K	2.12 [0.60; 10.33]	0.01 [-3.61; 3.31]	-0.50 [-3.96; 3.25]	0.33 [-3.62; 3.31]	-0.09 [-3.12; 3.70]	-0.40 [-3.32; 3.78]	0.24 [-2.99; 3.84]	-0.42 [-4.84; 3.11]	-0.80 [-5.05; 3.11]	-0.25 [-4.96; 3.09]
Phosphate binder med.	P-K	1.89 [1.25; 6.13]	1.01 [-2.11; 2.99]	-0.60 [-4.01; 3.17]	2.01 [-1.78; 5.02]	0.58 [-2.20; 2.63]	-0.34 [-4.75; 2.67]	2.14 [-2.67; 6.54]	0.85 [-1.18; 3.46]	-0.61 [-5.61; 4.33]	1.70 [-1.33; 4.36]
Blood fat medication	fl-Pi	-0.49 [-0.10; 0.73]	-0.16 [-2.09; 1.28]	-0.26 [-2.12; 1.30]	-0.07 [-2.08; 1.55]	-0.23 [-1.13; 0.54]	-0.44 [-1.27; 0.51]	-0.09 [-1.22; 0.67]	0.18 [-1.65; 1.80]	0.11 [-1.74; 1.82]	0.22 [-1.67; 1.98]
Protein, E%	fl-Pi	-0.27 [-0.26; 1.07]	0.06 [-0.87; 0.70]	-0.07 [-0.96; 0.70]	0.15 [-0.86; 0.83]	0.16 [-0.43; 1.15]	0.08 [-0.59; 1.19]	0.33 [-0.37; 1.21]	0.11 [-0.75; 1.38]	0.02 [-1.01; 1.49]	0.19 [-0.67; 1.46]
Calsium	P-K	0.04 [0.18; 2.18]	-0.17 [-1.88; 2.04]	-0.48 [-2.60; 2.34]	0.24 [-1.00; 2.30]	0.50 [-0.69; 1.75]	-0.16 [-1.44; 1.22]	0.84 [-0.63; 2.88]	0.26 [-0.83; 2.27]	-0.08 [-1.53; 1.82]	0.63 [-1.01; 2.61]
Phosphate binder med.	fl-Pi	-0.05 [-0.23; 1.67]	-0.18 [-1.40; 1.40]	-0.53 [-2.78; 1.76]	0.56 [-1.48; 1.86]	0.10 [-1.26; 1.40]	-0.35 [-1.78; 1.72]	0.58 [-1.56; 2.12]	-0.21 [-1.34; 1.04]	-0.36 [-1.53; 1.18]	0.04 [-1.88; 1.24]
Protein, g/kg	P-K	0.43 [-0.26; 4.73]	-0.93 [-5.97; 2.43]	-1.03 [-6.08; 2.43]	-0.85 [-5.87; 2.44]	-0.01 [-5.03; 2.51]	-0.18 [-5.04; 2.62]	0.14 [-4.87; 2.60]	0.42 [-2.78; 3.61]	0.29 [-2.77; 3.55]	0.54 [-2.82; 3.50]
Fat E%	fl-Pi	0.95 [0.68; 5.36]	0.36 [-2.04; 3.66]	0.12 [-2.17; 3.65]	0.67 [-2.21; 3.87]	0.55 [-1.09; 3.21]	0.33 [-1.34; 3.28]	0.79 [-1.12; 3.23]	0.06 [-2.57; 2.91]	-0.08 [-2.51; 3.07]	0.23 [-2.59; 3.16]
Monounsaturated Fatty Acids, E%	fl-Pi	-0.07 [0.19; 1.09]	-0.04 [-1.96; 1.01]	-0.15 [-2.07; 1.06]	-0.04 [-2.04; 1.05]	-0.14 [-1.25; 0.73]	-0.33 [-1.63; 0.76]	0.01 [-1.11; 0.97]	-0.08 [-1.04; 1.19]	0.01 [-1.01; 1.28]	0.03 [-1.02; 1.46]
Saturated Fatty Acids, E%	fl-Pi	-1.13 [-0.33; 1.61]	-0.15 [-1.88; 1.22]	-0.20 [-1.95; 1.16]	-0.12 [-1.82; 1.30]	-0.14 [-1.02; 1.12]	-0.19 [-1.10; 1.09]	-0.08 [-1.00; 1.24]	-0.02 [-1.20; 1.72]	-0.03 [-1.21; 1.64]	0.03 [-1.14; 2.00]
Calsium	fl-Pi	-0.01 [-0.39; 2.45]	-0.18 [-0.94; 0.74]	-0.26 [-0.91; 0.99]	0.00 [-0.94; 0.88]	-0.20 [-0.64; 0.20]	-0.41 [-0.98; 0.18]	0.01 [-0.63; 0.50]	-0.01 [-0.98; 1.08]	-0.04 [-1.09; 1.13]	0.18 [-0.95; 1.32]
Phosphorous	fl-Pi	0.02 [0.03; 2.03]	0.00 [-1.12; 1.54]	-0.12 [-1.35; 1.40]	0.09 [-1.01; 1.56]	0.03 [-1.00; 0.78]	-0.05 [-1.15; 0.81]	0.13 [-0.88; 0.97]	-0.12 [-1.60; 1.44]	-0.16 [-1.49; 1.37]	-0.09 [-1.56; 1.47]
Diabetes medication	fl-Pi	-0.24 [-0.31; 1.53]	-0.45 [-2.01; 1.01]	-0.66 [-2.46; 1.09]	-0.14 [-1.85; 1.16]	0.09 [-0.73; 1.09]	-0.17 [-1.06; 1.14]	0.22 [-0.87; 1.09]	0.64 [-1.26; 2.88]	0.60 [-1.32; 2.86]	0.85 [-1.13; 3.00]
Renavit	fl-Pi	-3.72 [-1.64; 2.92]	-1.90 [-10.81; 3.65]	-2.01 [-11.83; 3.68]	-1.70 [-9.90; 3.83]	-0.54 [-2.69; 2.10]	-0.66 [-2.90; 2.11]	-0.43 [-2.12; 2.23]	-2.77 [-14.84; 3.19]	-2.88 [-15.22; 3.15]	-2.71 [-14.52; 3.19]
Potassium	fl-Pi	-0.05 [-0.22; 0.72]	0.13 [-0.65; 1.08]	0.01 [-0.81; 1.01]	0.25 [-0.66; 1.58]	-0.07 [-0.58; 0.48]	-0.18 [-0.81; 0.47]	0.15 [-0.45; 1.47]	0.17 [-0.46; 0.90]	0.15 [-0.63; 1.06]	0.27 [-0.39; 1.37]
Energy, kcal/kg	fl-Pi	0.20 [0.39; 2.00]	0.30 [-1.74; 1.79]	0.21 [-1.83; 1.99]	0.43 [-1.63; 2.09]	0.63 [-1.09; 1.71]	0.53 [-1.25; 1.60]	0.77 [-0.97; 1.88]	0.48 [-2.37; 2.65]	0.42 [-2.55; 2.62]	0.57 [-2.27; 2.65]
Protein, g/kg	fl-Pi	-0.69 [-1.19; 2.13]	-0.55 [-1.93; 1.48]	-0.61 [-1.89; 1.52]	-0.49 [-1.84; 1.58]	-0.60 [-1.74; 1.19]	-0.75 [-2.09; 1.13]	-0.46 [-1.68; 1.19]	-0.77 [-2.30; 1.62]	-0.88 [-2.51; 1.56]	-0.68 [-2.41; 1.70]
Vitamin D	P-K	-0.04 [-0.15; 0.89]	-0.06 [-1.10; 0.72]	-0.14 [-1.18; 0.49]	0.22 [-0.85; 1.15]	-0.31 [-0.82; 0.19]	-0.49 [-1.61; 0.19]	-0.10 [-0.67; 0.57]	-0.06 [-0.73; 0.60]	-0.16 [-1.02; 0.53]	0.12 [-0.75; 0.80]
Fiber	P-K	0.46 [0.54; 2.53]	0.31 [-0.77; 1.04]	0.25 [-0.92; 1.07]	0.37 [-0.70; 1.08]	0.17 [-0.72; 1.26]	0.01 [-1.36; 0.94]	0.29 [-0.52; 1.84]	0.59 [-1.38; 1.93]	0.47 [-1.70; 1.81]	0.66 [-1.55; 2.12]
Potassium	P-K	0.02 [-0.15; 1.95]	-0.03 [-1.40; 1.53]	-0.25 [-1.91; 1.01]	0.01 [-1.34; 1.30]	0.17 [-0.91; 0.83]	-0.17 [-1.11; 0.55]	0.40 [-1.05; 1.45]	0.00 [-1.09; 2.00]	-0.10 [-1.17; 1.81]	0.20 [-1.08; 1.67]
Saturated Fatty Acids, E%	P-K	0.44 [0.61; 2.10]	0.09 [-2.07; 1.52]	-0.09 [-2.67; 1.49]	0.20 [-1.99; 1.60]	0.27 [-1.88; 1.67]	0.16 [-2.14; 1.74]	0.39 [-1.71; 1.80]	0.61 [-1.16; 1.99]	0.54 [-1.32; 2.00]	0.64 [-1.20; 2.08]
Water	fl-Pi	-0.08 [0.19; 0.89]	0.16 [-0.49; 0.81]	0.04 [-0.61; 0.85]	0.24 [-0.57; 1.22]	0.16 [-0.15; 0.58]	-0.05 [-0.54; 0.66]	0.28 [-0.23; 0.99]	0.12 [-1.01; 0.88]	0.04 [-1.21; 0.84]	0.26 [-1.00; 1.17]
Gender	fl-Pi	-0.10 [0.08; 1.79]	0.10 [-1.41; 1.42]	-0.05 [-1.73; 1.38]	0.19 [-1.40; 1.43]	-0.30 [-1.13; 0.78]	-0.50 [-2.01; 0.64]	-0.19 [-1.13; 1.14]	-0.58 [-1.89; 0.93]	-0.63 [-2.16; 1.00]	-0.51 [-2.02; 1.35]
Polyunsaturated Fatty Acids, E%	fl-Pi	-0.19 [-0.21; 0.62]	-0.23 [-1.36; 0.89]	-0.27 [-1.47; 0.90]	-0.15 [-1.24; 1.25]	-0.19 [-0.59; 0.35]	-0.26 [-0.87; 0.35]	-0.10 [-0.63; 0.60]	-0.08 [-0.88; 0.92]	-0.10 [-0.93; 0.95]	-0.04 [-0.88; 0.95]
Protein, E%	P-K	0.45 [0.74; 2.36]	0.39 [-1.22; 1.84]	0.34 [-1.25; 1.74]	0.52 [-1.10; 2.38]	-0.24 [-1.24; 1.60]	-0.37 [-1.46; 1.35]	-0.14 [-1.17; 2.11]	-0.36 [-1.92; 2.03]	-0.46 [-1.78; 1.80]	-0.27 [-1.89; 2.14]
Natrium	fl-Pi	0.20 [0.58; 1.47]	0.13 [-1.71; 1.14]	0.08 [-1.82; 1.15]	0.26 [-1.64; 1.44]	0.39 [-0.52; 1.13]	0.30 [-0.66; 0.93]	0.50 [-0.48; 1.39]	0.22 [-1.50; 1.89]	0.15 [-1.63; 1.74]	0.27 [-1.53; 2.08]
Salt	P-K	0.26 [0.58; 2.59]	1.90 [-0.98; 8.85]	1.68 [-0.90; 8.76]	2.14 [-0.92; 9.03]	0.42 [-0.61; 1.28]	0.19 [-0.73; 1.37]	0.72 [-0.51; 1.59]	0.81 [-2.20; 4.45]	0.57 [-2.43; 4.29]	1.04 [-2.17; 4.50]
Vitamin D	fl-Pi	-0.13 [0.02; 0.36]	-0.10 [-0.69; 0.32]	-0.17 [-0.81; 0.30]	0.08 [-0.62; 0.73]	0.08 [-0.23; 0.38]	-0.09 [-0.57; 0.43]	0.29 [-0.18; 0.67]	0.12 [-0.38; 0.71]	-0.03 [-0.51; 0.77]	0.22 [-0.50; 0.86]
Salt	fl-Pi	-0.08 [-0.42; 1.70]	-0.16 [-1.11; 1.58]	-0.30 [-1.15; 1.53]	-0.05 [-1.15; 1.57]	-0.34 [-0.96; 0.47]	-0.48 [-1.27; 0.50]	-0.15 [-0.89; 0.51]	-0.26 [-1.68; 1.38]	-0.42 [-1.70; 1.47]	-0.17 [-1.59; 1.52]

Overview of personal recommendations



Supplementary Figure S 5: Figure shows personal recommendations of potassium and phosphorous intake ($\hat{Q}^{min} - \hat{Q}^{max}$) as result of Algorithm (1) with predictive distributions of \hat{Q} in the background. In each row gives a numeric label of patient and type of patient's dialysis treatment (HD = hospital hemodialysis, HHD = home hemodialysis, PD = peritoneal dialysis). Personal recommendations are given for patients whose resulting concentrations levels could be predicted either over 90% or 80% probability. For the patients that did not receive confident enough personal recommendation reasoning of failure is indicated. Three right-most panels show the estimated concentrations matching the recommended intake. In these panels, grey bars indicate the estimated concentration without the effect of potassium and phosphorous intake, μ_{q_0} . All blue bars indicate the range of concentration that is resulted over the required $P^{max} > 80\%$ confidence by modifying potassium and phosphorous intake. The best exact probabilities are shown with percentage figures. Red bars indicate that even best concentration estimation has $P^{max} < 80\%$ confidence and personal recommendations are not given for these patients. For all the concentrations a maximum reached confidence P_m^{max} is reported, except for patients 22,23, and 29 only maximum overall confidence P^{max} is reported. The figure is plotted with ggplot2 package for R language (v 3.3.5, <https://ggplot2.tidyverse.org>).

Personal recommendation for a patient



Supplementary Figure S 6: Figure shows detailed intake recommendation for patient 10 from the previous recommendation table. The intake plot on the left shows the posterior samples of diet configurations that result concentrations of plasma potassium (P-K), fasting plasma phosphate (fP-Pi) and plasma albumin (P-Alb) to stay within their recommended limits. These levels are marked with vertical solid lines in the concentration panels. The black point in the middle of intake plot represents the patient's current potassium and phosphorous intake. Current concentrations matching this intake are shown with dashed vertical lines. Reported recommendation is shown with a rectangle that contains 95% of diet proposals that result recommended concentrations over 90% accuracy. Colouring of the rectangle sides match the concentration estimates in the right hand panels. The figure is plotted with ggplot2 package for R language (v 3.3.5, <https://ggplot2.tidyverse.org>).