PhUSE GPP Steering Board

Good Programming Practice at a Glance

The GPP Steering Board is a voluntary industry group with representatives from a diverse array of health and life sciences organizations

- Working to develop common good programming practices applicable to statistical programming for the development of high quality and efficient programming code
- Focus on analysis and reporting of clinical trial data; data integration; preparation of clinical data for e-submissions
- This Poster contains an overview of PhUSE GPP guidance Version 1: Consolidated industry guideline with potential to replace/augment individual company guidance and serve as a set of business rules when working on standard scripts for public use.



PhUSE GPP Guidance is found on PhUSE Wiki

The Wiki is a place to share your ideas and post your contributions.





http://www.phusewiki.org/wiki/index.php?title=Good Programming Practice Guidance

Before you begin

Understand what you need to do; plan how to get there; collect what you need

- SOPs & training
- Study Protocol
- Standards
- **Annotated CRF**
- SAP & Shells
- Specifications
- Tools and macros

GPP Principles

Write code that is

- Clear and easy to read and review
- Easy to maintain, modify and debug
- Robust and needs minimal code maintenance
- Can be reused or easily adapted.
- Efficient and uses minimal computer processing resource
- Written in such a way to reduce logical and syntax errors

Coding conventions

- Do not overwrite existing datasets
- Use of all uppercase should be avoided
- Separate data steps and procedures with at least one blank line – Use whitespace
- Use 'data=dataset' option in procedure statements
- End data steps and procedures with run or quit
- Put each statement on a separate line
- Indent statements belonging to a level by 2 to 5 columns (use the same number of spaces throughout the program), i.e. every nesting level should be visibly indented from the previous level.
- Do loops and if then else statements start and end with same level of indentation

Have a look at the Wiki, there is more ...

Examples of Good and bad code Review the 2 code snippets below: Both produce the same output. Why is code on the right better?

```
|data alb tmp ;
set &derived target..alb;
if saffl = 'Y';
if aval is missing then do;
tempflag1='x';
tempflag2='y';
end;
|data alb tmp( where = (avisitn is not missing));
 set alb tmp;
where (aval is not missing or chg is not missing);
run;
|proc sql;
select armcd, count(distinct subjid) into :arm1 - :arm2 , :trt1 - :trt2
from &derived target..asl
where saffl = 'Y'
group by armcd;
create table comb_alb as
select subjid, avisit, avisitn, "a" as groupby, paramcd, armcd, aval as val from
alb_tmp
where aval ne .
union all
select subjid, avisit, avisitn, "b" as groupby,
paramed, armed, chg as val from alb tmp
where chg ne . ;
quit;
* Sort the data ;
proc sort
data = comb alb;
by groupby paramed armed avisitn subjid;
run;
```

- (KEEP=studyid subjid usubjid avisit avisitn paramcd aval chg armcd tempflag); IF avisitn is missing then put "Note: Missing avisitn for " usubjid= paramcd= aval= chg= WHERE saffl = 'Y' GROUP BY armcd; ***************** DATA alb nomiss; WHERE (aval IS NOT MISSING or chg IS NOT MISSING) AND (avisith IS NOT MISSING); PROC SQL NOPRINT, CREATE TABLE comb alb AS SELECT subjid, avisit, avisitn, "Actual" AS groupby , paramcd, armcd, aval AS val FROM alb nomiss WHERE aval NE SELECT subjid, avisit, avisitn, "Change_baseline" AS groupby , paramed, armed, chg AS val FROM alb nomiss WHERE chg NE .; PROC SORT DATA = comb alb OUT = sort Alb; BY groupby paramed armed avisitn subjid;
- Comments explain the purpose of what is being done
- Indented Code
- Keep only the variables you need
- Data sets not overwritten
- Where clauses do not overwrite
- What else?

Layout & Header

- Always have a program header!
- Clean up work area from previous program
- Read in external data
- For multiple data sources, pre-process, then combine before common derivations are applied
- Process and manipulate data, merge and derive variables
- Perform analysis
- Report data: Produce tables, figures and listings
- Clean up work area

PROGRAM PROGRAM LOCATION	: xxxx.xx	CREATION DATE: YYYY-MM-DD
	: xxxx.sas	
	: xxxx.sas	
PROGRAM LOCATION		
PROGRAM LOCATION		
	: YYYY/YYYY/Y x Y	
PROGRAMMER	:	
DESCRIPTION	: <describe pu<="" td="" the=""><td>rpose of the program></td></describe>	rpose of the program>
FINAL PROGRAM DATE	: < Date the progr	am was considered final/validated>
SOFTWARE VERSION	: <state n<="" sotware="" td=""><td>ame, version, operating system></td></state>	ame, version, operating system>
REVISION HISTORY (COE	Y FOR EACH REVISION) :
DATE	:	
PROGRAMMER		
REASON FOR CHANGE	:	

DO

- Use a program Header
- Use Informative comments in appropriate places
- Use unique and
- informative dataset names Use one statement per

missing and unexpected

Indent consistently Program defensively for

data

Check log for ERRORS, WARNING, and NOTES that indicate data issues

DON'T

- Use a header but forget to fill in or update!
- Use comments that simply say what the code does (not Why)
- Use inconsistent or random naming for datasets and variables
- Program around data issues or "to the data"
- Ignore Notes in the Log that indicate potential issues E.G. Automatic Number/character conversions, Merge by multiple records

Check your Log!

Not(e): just ERROR:s and WARNING:s! not referenced never been refere not resolved is not in the report def has more than one data set with repeats of by values current word or quoted string has become more than 200 observation(s) outside the axis range is unknown uninitialized cannot be determined extraneous information missing values were generated observation(s) contained a MISSING value Invalid arguments truncated to 32 characters overwritten by can't modify it at this time have been converted to

GPP Steering Board		
Mark Foxwell, Chair	PRA Health Sciences UK	
Beate Hientzsch	Accovion, Germany	
Shafi Chowdhury	Shafi Consultancy, UK	
Dean Grundy	Roche, UK	
Jennifer Chin	Eisai, UK	
Cindy Song	Sanofi, US	
Kate Peacock	Quintiles, UK	
Alexandra Marquat	Boehringer-Ingelheim Germany	
Maria Dalton	GSK, US	
Gayathri Kolandaivelu	Jannsen R&D (J&J), US	
Jane Marrer	Merck, US	
Ninan Luke	Novartis, India	
Art Collins	Biogen Idec, US	
Ralf Gessenich	Grunenthal, Germany	