

Alain Smits

Phuse SDE Belgium

SUPERCHARGING PROC CONTENTS

Introduction

- ⦿ Combines a structure and contents overview in a dynamic web page
- ⦿ Easier to explore new data sets, especially complex ones
- ⦿ Pick up data issues quickly

The macro

```
%dscont(inds      = ,  
          limit    = ,  
          path     = ,  
          file     = );
```

Example

```
%dscont(inds = sds.1b);
```

Default output to work folder, file opens
automatically in SAS DMS

Sample output

ADAM.LBAD

Laboratory Results Analysis Dataset

Created: 19OCT10:15:50:19

Obs : 9750

Key :

#	VARIABLE	LABEL	TYPE	LENGTH	+	CONTENTS
1	STUDYID	STUDY IDENTIFIER	CHAR	40	+	1 distinct values, 0 missing values
2	SITEID	STUDY SITE IDENTIFIER	CHAR	10	+	11 distinct values, 0 missing values
3	INVMAM	INVESTIGATOR NAME	CHAR	120	+	11 distinct values, 0 missing values
4	SUBJID	SUBJECT IDENTIFIER FOR THE STUDY	CHAR	20	+	27 distinct values, 0 missing values
5	USUBJID	UNIQUE SUBJECT IDENTIFIER	CHAR	40	+	27 distinct values, 0 missing values
6	SUBJINIT	SUBJECT INITIALS	CHAR	5	+	27 distinct values, 0 missing values
7	AGE	AGE (years) AT REFERENCE DATE/TIME	NUM	8	+	3 distinct values, 0 missing values Mean: 3.968 Minimum: 3 Maximum: 5
8	SEX	SEX	CHAR	1	+	2 distinct values, 0 missing values
9	RACE	RACE	CHAR	80	+	4 distinct values, 0 missing values
10	ARMCD	PLANNED ARM CODE	CHAR	8	+	1 distinct values, 0 missing values
11	ARM	DESCRIPTION OF PLANNED ARM	CHAR	200	+	1 distinct values, 0 missing values
12	ARMN	PLANNED ARM NUM. CODE	NUM	8	+	1 distinct values, 0 missing values Mean: 1 Minimum: 1 Maximum: 1 Mis
13	LBGRPID	GROUP ID	CHAR	120	+	214 distinct values, 227 missing values
14	PHASE	PHASE	CHAR	80	+	3 distinct values, 0 missing values
15	PHASEN	PHASE (NUM CODE)	NUM	8	+	3 distinct values, 0 missing values Mean: 0.864 Minimum: 0 Maximum: 2
16	VISITNUM	VISIT NUMBER	NUM	8	+	16 distinct values, 0 missing values Mean: 17.618 Minimum: 1 Maximum: 1
17	VISIT	VISIT NAME	CHAR	120	+	16 distinct values, 0 missing values
18	ANALTPT	ANALYSIS TIMEPOINT	CHAR	100	+	10 distinct values, 0 missing values

Sample output

44	LBSPEC	SPECIMEN TYPE	CHAR	40	+	4 distinct values, 1760 missing values
45	LBFAST	FASTING STATUS	CHAR	1	+	3 distinct values, 227 missing values
46	LBTOX	TOXICITY	CHAR	40	+	9 distinct values, 440 missing values
47	LBTOXN	TOXICITY NUMBER	NUM	8	+	9 distinct values, 440 missing values Mean: 2.61 Minimum: 0 Maximum: 7 Missing
48	LBTOXNM	TOXICITY NAME	CHAR	60	-	49 distinct values, 385 missing values Only approx. 30 values shown 385x **MISSING** 224x ALANINE AMINO TRANSFERASE 225x ALKALINE PHOSPHATASE 135x ALPHA-1 ACID GLYCOPROTEIN 219x AMYLASE 218x ASPARTATE AMINO TRANSFERASE 174x BASOPHILS 221x BASOPHILS (%) 225x BICARBONATE 218x BLOOD UREA NITROGEN 227x CALCIUM 223x CHLORIDE 227x CREATININE 219x DIRECT BILIRUBIN 174x EOSINOPHILS 219x EOSINOPHILS (%) 220x HAEMATOCRIT 221x HAEMOGLOBIN 128x HIGH DENSITY LIPOPROTEIN 225x HYPERBILIRUBINEMIA

Under the hood

⦿ Programming challenges

- Performance when processing large data sets
- Handling user defined formats
- Avoiding information overload in the output

Under the hood

⦿ Performance

- Proc tabulate provides incredible processing efficiency to create statistics on all variables.
- Requires a fair amount of data tweaking to get a useful data format for further processing due to the special format of the output data set of proc tabulate

Under the hood

⦿ User defined formats

- The macro needs to collect the user defined formats to ensure the table correctly displays the values, although the format itself is currently not displayed
- In our organization, the use of user defined formats is limited
- Code can be simplified through the recently introduced `vvalue()` function that returns the formatted value of a variable

Under the hood

- ◎ Avoiding information overload
 - The use of dynamic HTML (JavaScript) allows information to be collapsed to allow a clean structure view and expanded for a contents overview

Extras

- ⦿ Saving as an excel file
 - Useful for copy and pasting.

Future enhancements

- (User Defined) Formats are currently not shown because they are not used in SDTM or ADaM.



Contact details

Alain Smits
Associate Director Clinical Programming
Janssen R&D
Beerse, Belgium
+32 14 64 1325 or +32 495 26 90 08
asmits@its.jnj.com