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# Translation from narrative text to standard codes variables with Stata

Federico Belotti University of Rome "Tor Vergata" Rome, Italy federico.belotti@uniroma2.it Domenico Depalo
Bank of Italy
Rome, Italy
domenico.depalo@bancaditalia.it

**Abstract.** In this article, we describe **screening**, a new Stata command for data management that can be used to examine the content of complex narrative-text variables to identify one or more user-defined keywords. The command is useful when dealing with string data contaminated with abbreviations, typos, or mistakes. A rich set of options allows a direct translation from the original narrative string to a user-defined standard coding scheme. Moreover, **screening** is flexible enough to facilitate the merging of information from different sources and to extract or reorganize the content of string variables.

**Editors' note.** This article refers to undocumented functions of Mata, meaning that there are no corresponding manual entries. Documentation for these functions is available only as help files; see help regex.

**Keywords:** dm0050, screening, keyword matching, narrative-text variables, standard coding schemes

### 1 Introduction

Many researchers in varied fields frequently deal with data collected as narrative text, which are almost useless unless treated. For example,

- Electronic patient records (EPRs) are useful for decision making and clinical research only if patient data that are currently documented as narrative text are coded in standard form (Moorman et al. 1994).
- When different sources of data use different spellings to identify the same unit of interest, the information can be exploited only if codes are made uniform (Raciborski 2008).
- Because of verbatim responses to open-ended questions, survey data items must be converted into nominal categories with a fixed coding frame to be useful for applied research.

These are only three of the many critical examples that motivate an ad hoc command.

Recoding a narrative-text variable into a user-defined standard coding scheme is currently possible in Stata by combining standard data-management commands (for example, generate and replace) with regular expression functions (for example, regexm()).

However, many problems do not yield easily to this approach, especially problems containing complex narrative-text data. Consider, for example, the case when many source variables can be used to identify a set of keywords; or the case when, looking at different keywords, one is within a given source variable but not necessarily at the beginning of that variable, whereas the others are at the beginning, the end, or within that or other source variables. Because no command jointly handles all possible cases, these cases can be treated with existing Stata commands only after long and tedious programming, increasing the possibility of introducing errors. We developed the screening command to fill this gap, simplifying data-cleaning operations while being flexible enough to cover a wide range of situations.

In particular, screening checks the content of one or more string variables (sources) to identify one or more user-defined regular expressions (keywords). Because string variables are not flexible, to make the command easier and more useful, a set of options reduces your preparatory burden. You can make the matching task wholly case insensitive or set matching rules aimed at matching keywords at the beginning, the end, or within one or more sources. If source variables contain periods, commas, dashes, double blanks, ampersands, parentheses, etc., it is possible to perform the matching by removing such undesirable content. Moreover, if the matching task becomes more difficult because of abbreviations or even pure mistakes, screening allows you to specify the number of letters to screen in a keyword. Finally, the command allows a direct translation of the original string variables in a user-defined standard coding scheme.

All these features make the command simple, extremely flexible, and fast, minimizing the possibility of introducing errors. It is worth emphasizing that we find Mata more convenient to use than Stata, with advantages in terms of time execution.

The article is organized as follows. In section 2, we describe the new screening command, and we provide some useful tips in section 3. Section 4 illustrates the main features of the command using EPR data, while section 5 details some critical cases in which the use of screening may aid your decision to merge data from different sources or to extract and reorder messy data. In the last section, section 6, we offer a short summary.

# 2 The screening command

String variables are useful in many practical circumstances. A drawback is that they are not so flexible: for example, in EPR data, coding CHOLESTEROL is different from coding CHOLESTEROL LDL, although the broad pathology is the same. Stata and Mata offer many built-in functions to handle strings. In particular, screening extensively uses the Mata regular-expression functions regexm(), regexr(), and regexs().

(Continued on next page)

#### 2.1 Syntax

```
screening [if] [in], sources(varlist[, sourcesopts]) keys([matching_rule] "string" [...]) [letters(#) explore(type) cases(newvar) newcode(newvar [, newcodeopts]) recode(recoding_rule "user_defined_code" [recoding_rule "user_defined_code" [...]) checksources tabcheck memcheck nowarnings save time]
```

#### 2.2 Options

sources(varlist[, sourcesopts]) specifies one or more string source variables to be screened. sources() is required.

sources opts	description
lower	perform a case-insensitive match (lowercase)
upper	perform a case-insensitive match (uppercase)
trim	match keywords by removing leading and trailing blanks
	from sources
itrim	match keywords by collapsing sources with consecutive
	internal blanks to one blank
removeblank	match keywords by removing from sources all blanks
removesign	match keywords by removing from sources the following
	signs: * + ? / \ % ( ) [ ] { }   . ^ # \$

keys([matching\_rule] "string" ...) specifies one or more regular expressions (keywords) to be matched with source variables. keys() is required.

$matching\_rule$	description
begin end	match keywords at beginning of string match keywords at end of string

letters (#) specifies the number of letters to be matched in a keyword. The number of letters can play a critical role: specifying a high number of letters may cause the number of matched observations to be artificially low because of mistakes or abbreviations in the source variables; on the other hand, matching a small number of letters may cause the number of matched observations to be artificially high because of the inclusion of uninteresting cases containing the "too short" keyword. The default is to match keywords as a whole.

explore(type) allows you to explore screening results.

type	description
tab count	tabulate all matched cases for each keyword within each source variable display a table of frequency counts of all matched cases for each keyword within each source variable

cases (newvar) generates a set of categorical variables (as many as the number of keywords) showing the number of occurrences of each keyword within all specified source variables.

newcode(newvar[, newcodeopts]) generates a new (numeric) variable that contains the position of the keywords or the regular expressions in keys(). The coding process is driven by the order of keywords or regular expressions.

newcode opts	description
replace	replace newvar if it already exists
add	obtain <i>newvar</i> as a concatenation of subexpressions returned by
	regexs(n), which must be specified as a
	$user\_defined\_code$ in recode
label	attach keywords as value labels to newvar
numeric	convert <i>newvar</i> from string to numeric; it can be specified only if the <b>recode()</b> option is specified

recode(recoding\_rule "user\_defined\_code" [recoding\_rule "user\_defined\_code" ...])
recodes the newcode() newvar according to a user-defined coding scheme. recode()
must contain at least one recoding\_rule followed by one user\_defined\_code. When you
specify recode(1 "user\_defined\_code"), the "user\_defined\_code" will be used to recode all matched cases from the first keyword within the list specified via the keys()
option. If recode(2,3 "user\_defined\_code") is specified, the "user\_defined\_code" will
be used to recode all cases for which second and third keywords are simultaneously
matched, and so on. This option can only be specified if the newcode() option is
specified.

checksources checks whether source variables contain special characters. If a matching rule is specified (begin or end via keys()), checksources checks the sources' boundaries accordingly.

tabcheck tabulates all cases from checksources. If there are too many cases, the option does not produce a table.

memcheck performs a "preventive" memory check. When memcheck is specified, the command will exit promptly if the allocated memory is insufficient to run screening. When memory is insufficient and screening is run without memcheck, the command could run for several minutes or even hours before producing the message no room to add more variables.

nowarnings suppresses all warning messages.

save saves in r() the number of cases detected, matching each source with each keyword.

time reports elapsed time for execution (seconds).

# 3 Tips

The low flexibility of string variables is a reason for concern. In this section, we provide some tips to enhance the usefulness of **screening**. Some tips are useful to execute the command, while other tips are useful to check the results.

Most importantly, capitalization matters: this means that screening for KEYWORD is different from screening for keyword. If source variables contain HEMINGWAY and you are searching for Hemingway, screening will not identify such keyword. If suboption upper (lower) is specified in sources(), keywords will be automatically matched in uppercase (lowercase).

Choose an appropriate *matching\_rule*. The screening default is to match keywords over the entire content of source variables. By specifying the *matching\_rule* begin or end within the keys() option, you may switch accordingly the matching rule on string boundaries. For example, if sources contain HEMINGWAY ERNEST and ERNEST HEMINGWAY and you are searching begin HEMINGWAY, the screening command will identify the keyword only in the former case. Whether the two cases are equivalent must be evaluated case by case.

Another issue is how to choose the optimal number of letters to be screened. For example, with EPR data, different physicians might use different abbreviations for the same pathologies. And so talking about a "right" number of letters is nonsense. As a rule of thumb, the number of letters should be specified as the minimum number that uniquely identifies the case of interest. Using many letters can be too exclusive, while using few letters can be too inclusive. In all cases, but in particular when the appropriate number of letters is unknown, we find it useful to tabulate all matched cases through the explore(tab) option. Because it tabulates all possible matches between all keywords and all source variables, it is the fastest way to explore the data and choose the best matching strategy (in terms of keywords, matching rule, and letters).

Advanced users can maximize the potentiality of screening by mixing keywords with Stata regular-expression operators. Mixing in operators allows you to match more-complex patterns, as we show later in the article. For more details on regular-expression syntaxes and operators, see the official documentation at http://www.stata.com/support/faqs/data/regex.html.

<sup>1.</sup> The letters() option does not work if a keyword contains regular-expression operators.

screening displays several messages to inform you about the effects of the specified options. For example, consider the case in which you are searching some keywords containing regular-expression operators. screening will display a message with the correct syntax to search a keyword containing regular-expression operators. The nowarnings option allows you to suppress all warning messages.

screening generates several temporary variables (proportional to the number of keywords you are looking for and to the number of sources you are looking from). So when you are working with a big dataset and your computer is limited in terms of RAM, it might be a good idea to perform a "preventive" memory check. When the memcheck option is specified and the allocated memory is insufficient, screening will exit promptly rather than running for several minutes or even hours before producing the message no room to add more variables.

We conclude this section with an evaluation of the command in terms of time execution using different Stata flavors and different operating systems. In particular, we compare the latest version of screening written using Mata regular-expression functions with its beta version written entirely using the Stata counterpart. We built three datasets of 500,000 (A), 5 million (B), and 50 million (C) observations with an ad hoc source variable containing 10 different words: HEMINGWAY, FITZGERALD, DOSTOEVSKIJ, TOLSTOJ, SAINT-EXUPERY, HUGO, CERVANTES, BUKOWSKI, DUMAS, and DESSI. Screening for HEMINGWAY (50% of total cases) gives the following results (in seconds):

Stata flavor and		Mata	a		Stat	 a
operating system	A	В	$\mathbf{C}$	A	В	$\mathbf{C}$
Stata/SE 10 (32-bit) and Mac os X 10.5.8 (64-bit)*	0.66	6.67	na	0.93	9.24	na
Stata/MP 11 (64-bit) and Mac os X 10.5.8 (64-bit)*	0.60	5.66	na	0.85	7.73	na
Stata/MP 11 (64-bit) and Window Server 2003 (64-bit) $^+$	0.37	3.70	37.22	0.70	7.06	70.59

<sup>\*</sup> Intel Core 2 Duo 2.2 GHz (dual core) with 4 GB RAM

The table speaks for itself!

# 4 Example

To illustrate the command, we use anonymized patient-level data from the Health Search database, a nationally representative panel of patients run by the Italian College of General Practitioners (Italian Society of General Medicine). Our sample contains freely inputted EPRs concerning the prescription of diagnostic tests.<sup>2</sup> A list of 15 observations

<sup>&</sup>lt;sup>+</sup> AMD Opteron 2.2 GHz (quad core) with 20 GB RAM

<sup>2.</sup> The original data are in Italian. Where necessary for comprehension, we translate to English.

from the "uppercase" source variable diagn\_test\_description provides an overview of cases at hand:

. list diagn\_test\_descr in 1/15, noobs separator(20)

diagn\_test\_descr TRIGLICERIDI EMOCROMO FORMULA COLESTEROLO TOTALE ALTEZZA PT TEMPO PROTROMBINA VISITA CARDIOLOGICA CONTROLLO HCV AB EPATITE C COMPONENTE MONOCLONALE ATTIVITA' FISICA PSA ANTIGENE PROSTATICO SPECIFICO RX CAVIGLIA SN FAMILIARITA' K UTERO TRIGLICERIDI URINE ESAME COMPLETO URINE PESO SPECIFICO

As you can see, this is a rich EPR dataset that is totally useless unless treated. If data were collected for research purposes, physicians would be given a finite number of possible options. There is much agreement in the scientific community that the cost to leave the burden of inputting standard codes directly to physicians at the time of contact with the patient is higher than the relative benefit: the task is extremely onerous, it is unrelated to the physician's primary job, and most importantly, it requires extra effort. Therefore, the common view supports the implementation of data-entry methods that do not disturb the physician's workflow (Yamazaki and Satomura 2000).

From the above list of observations, it is also clear that free-text data entry provides physicians with the freedom to determine the order and detail at which they want to input data. Even if the original free-text data were complete, it would still be difficult to extract standardized and structured data from this kind of record because of abbreviations, typos, or mistakes (Moorman et al. 1994). Extracting data in the presence of abbreviations and typos is exactly what screening allows you to do.

As a practical example, we focus on the identification of different types of cholesterol tests. In particular, our aim is to create a new variable (diagn\_test\_code) containing cholesterol test codes according to the Italian National Health System coding scheme. Because at least three types of cholesterol test exist, namely, hdl, ldl, and total, our matching strategy must take into account that a physician can input 1) only the types of the test, 2) only its broad definition (cholesterol), or 3) both, without considering abbreviations, typos, mistakes, and further details.

Thus we first explore the data by running screening with the explore(tab) option:

. screening, sources(diagn\_test\_descr, lower) keys(colesterolo) explore(tab) Cases of colesterolo found in diagn\_test\_descr

colesterolo	Freq.	Percent	Cum.
colesterolo totale	2,954	51.86	51.86
hdl colesterolo	1,854	32.55	84.41
ldl colesterolo	617	10.83	95.24
colesterolo hdl	117	2.05	97.30
colesterolo ldl	37	0.65	97.95
colesterolo tot	28	0.49	98.44
colesterolo	24	0.42	98.86
colesterolo hdl sangue	16	0.28	99.14
colesterolo totale sangue	16	0.28	99.42
colesterolo esterificato	4	0.07	99.49
colesterolo tot.	4	0.07	99.56
colesterolo hdl 90.14.1	3	0.05	99.61
colesterolo totale 90143	3	0.05	99.67
colesterolo libero	2	0.04	99.70
colesterolo stick	2	0.04	99.74
colesterolo tot hdl	2	0.04	99.77
colesterolo totale 90.143	2	0.04	99.81
ultima misurazione colesterolo	2	0.04	99.84
colesterolo hdl	1	0.02	99.86
colesterolo ldl 90.14.2	1	0.02	99.88
colesterolo non ldl	1	0.02	99.89
colesterolo t. mg/dl	1	0.02	99.91
colesterolo tot. c	1	0.02	99.93
colesterolo tot. hdl	1	0.02	99.95
colesterolo tot.,	1	0.02	99.96
colesterolo totale h	1	0.02	99.98
rich, specialistica colesterolo trigl	1	0.02	100.00
Total	5,696	100.00	

Here the lower suboption makes the matching task case insensitive. Apart from the explore(tab) option, the syntax above is compulsory and performs what we call a default matching, that is, an exact match of the keyword colesterolo over the entire content of the source variable diagn\_test\_descr. The tabulation above (notice the lowercase) informs you that the keyword colesterolo is encountered in 5,696 cases. What do these cases contain? Because you did not instruct the command to match a shorter length of the keyword, the only possible case is the keyword itself; all the cases contain the keyword colesterolo.

Given the nature of the data, it might be convenient to run screening with a shorter length of the keyword so as to find possible partial matching in the presence of abbreviations or mistakes. The letters(#) option instructs screening to perform the match on a shorter length:

(Continued on next page)

. screening, sources(diagn\_test\_descr, lower) keys(colesterolo) letters(5)
> explore(tab)

Cases of coles found in diagn\_test\_descr

coles	Freq.	Percent	Cum.
colesterolo totale	2,954	37.25	37.25
hdl colesterolo	1,854	23.38	60.62
coles ldl	1,343	16.93	77.56
hdl colest	853	10.76	88.31
ldl colesterolo	617	7.78	96.09
colesterolo hdl	117	1.48	97.57
colesterolo ldl	37	0.47	98.03
colesterolo tot	28	0.35	98.39
colesterolo	24	0.30	98.69
colesterolo hdl sangue	16	0.20	98.89
colesterolo totale sangue	16	0.20	99.09
colesterolemia	14	0.18	99.27
hdl colest.	5	0.06	99.33
colest.tot.	4	0.05	99.38
colesterolo esterificato	4	0.05	99.43
colesterolo tot.	4	0.05	99.48
azotemia glicemia colest	3	0.04	99.52
colest. hdl	3	0.04	99.56
colesterolo hdl 90.14.1	3	0.04	99.60
colesterolo totale 90143	3	0.04	99.63
colesterolo libero	2	0.03	99.66
colesterolo stick	2	0.03	99.68
colesterolo tot hdl	2	0.03	99.71
colesterolo totale 90.143	2	0.03	99.74
ldl colest.	2	0.03	99.76
ultima misurazione colesterolo	2	0.03	99.79
colest. ldl	1	0.01	99.80
colest. tot.	1	0.01	99.81
colest.tot	1	0.01	99.82
colester.tot.hdl,	1	0.01	99.84
colesterolo hdl	1	0.01	99.85
colesterolo ldl 90.14.2	1	0.01	99.86
colesterolo non ldl	1 1	0.01 0.01	99.87
colesterolo t. mg/dl colesterolo tot. c			99.89
	1	0.01	99.90
colesterolo tot. hdl	1	0.01	99.91
colesterolo tot.,	1	0.01	99.92
colesterolo totale h	1	0.01	99.94
emocromo c. colester	1 1	0.01 0.01	99.95
glicemia colesterolemia-	1	0.01	99.96
<pre>got gpt colest / trigli/creat/emocromo   rich, specialistica colesterolo trigl</pre>	1	0.01	99.97 99.99
uricemia uricuria colest	1	0.01	100.00
uricemia uricuria corest	1	0.01	
Total	7,931	100.00	

By specifying a five-letter partial match, screening detects 2,235 new cases of cholesterol tests. By further reducing the number of letters, we get the following result:<sup>3</sup>

<sup>3.</sup> Because of space restrictions, we deliberately omit the complete tabulation obtainable with the explore(tab) option. It is available upon request.

. screening, sources(diagn\_test\_descr, lower) keys("colesterolo") letters(3)
> explore(tab)

Cases of col found in diagn\_test\_descr

col	Freq.	Percent	Cum.
colesterolo totale	2,954	23.45	23.45
col tot	2,034	16.15	39.60
hdl colesterolo	1,854	14.72	54.32
coles ldl	1,343	10.66	64.99
hdl colest	853	6.77	71.76
ldl colesterolo	617	4.90	76.66
urinocoltura coltura urina	326	2.59	79.25
v.ginecologica	161	1.28	80.52
eco tiroide eco capo e collo	150	1.19	81.71
colesterolo hdl	117	0.93	82.64
(output omitted)			
colesterolo ldl	37	0.29	90.77
calcolo rischio cardiovascolare (iss)	35	0.28	91.04
coprocoltura coltura feci	33	0.26	91.31
colore	32	0.25	91.56
ecocolordoppler arti inf. art.	32	0.25	91.81
urinocoltura	32	0.25	92.07
colposcopia	31	0.25	92.31
colesterolo tot	28	0.22	92.54
reticolociti	28	0.22	92.76
ecodoppler a.inferiori ecocolor venosa	27	0.21	92.97
eco ginecologica	25	0.20	93.17
colesterolo	24	0.19	93.36
rischio cardio vascolare nota 13	23	0.18	93.55
rischio cardiovascolare % a 10 anni	22	0.17	93.72
ecodoppler a.inferiori ecocolor arter.	19	0.15	93.87
(output omitted)			
col hdl	3	0.02	97.13
colest. hdl	3	0.02	97.16
colesterolo hdl 90.14.1	3	0.02	97.18
colesterolo totale 90143	3	0.02	97.21
conta batt.,urinocoltura, antibiogramma	3	0.02	97.23
eco cardiaca con doppler e colordoppler	3	0.02	97.25
eco color/doppl.car. ver	3	0.02	97.28
eco(color)dopplergrafia	3	0.02	97.30
ecocardiografia colordoppler	3	0.02	97.32
ecocolordoppler art.aa.inf.	3	0.02	97.35
ecocolordoppler arterioso arti inferior	3	0.02	97.37
ecocolordoppler tronchi sovraortici	3	0.02	97.40
ecocolordopplergrafia cardiaca	3	0.02	97.42
ecografia muscolotendinea	3	0.02	97.44
ecografia tiroide eco capo e collo	3	0.02	97.47
familiarita ev.cerebrovascol.( 72m 74f	3	0.02	97.49
immunocomplessi circolanti	3	0.02	97.51
rx digerente (tenue e colon)	3	0.02	97.54
test broncodilatazione farmacologica	3	0.02	97.56
test cardiovascolare da sforzo con cicl	3	0.02	97.59
test sforzo cardiovascol. pedana mobile	3	0.02	97.61
urinocoltura atb+mic	3	0.02	97.63
urinocoltura con antibiogramma	3	0.02	97.66
urinocoltura identificazione batt.+ ab	3	0.02	97.68
che colinesterasi	2	0.02	97.70
col	2	0.02	97.71

colangio rm	2	0.02	97.73
colesterolo libero	2	0.02	97.75
colesterolo stick	2	0.02	97.76
colesterolo tot hdl	2	0.02	97.78
colesterolo totale 90.143	2	0.02	97.79
(output omitted)	'		
ldl colest.	2	0.02	98.11
(output omitted)	•		
col tot 216 hdl 58 fibri	1	0.01	98.48
col=245ldl=193tr=91	1	0.01	98.48
colangiografia intravenosa	1	0.01	98.49
colecistografia	1	0.01	98.50
colecistografia per os c	1	0.01	98.51
colest. ldl	1	0.01	98.52
colest. tot.	1	0.01	98.52
colest.tot	1	0.01	98.53
colester.tot.hdl,	1	0.01	98.54
colesterolo hdl	1	0.01	98.55
colesterolo ldl 90.14.2	1	0.01	98.55
colesterolo non ldl	1	0.01	98.56
colesterolo t. mg/dl	1	0.01	98.57
colesterolo tot. c	1	0.01	98.58
colesterolo tot. hdl	1	0.01	98.59
colesterolo tot.,	1	0.01	98.59
colesterolo totale h	1	0.01	98.60
colloquio psicologico	1	0.01	98.61
(output omitted)	•		
hdl col	1	0.01	99.22
(output omitted)			
visita specialistica colonscopia con bi	1	0.01	99.99
yersinia coltura feci	1	0.01	100.00
Total	12,595	100.00	

Again screening detects new cases: 2,034 cases characterized by the abbreviation col tot (that is, total cholesterol) that are impossible to identify without further reducing the number of letters. The problem is that, among all matched cases (12,595), there are also a number of unwanted cases, that is, cases containing the same spelling of the keyword but related to another type of diagnostic test. Despite this incorrect identification, we will show later in the section how to obtain a new "recoded variable" by specifying the appropriate recoding\_rule as an argument of the recode() option.

The number of letters you match plays a critical role: specifying a high number of letters may cause the number of matched observations to be artificially low due to mistakes or abbreviations in the source variables; on the other hand, matching a small number of letters may cause the number of matched observations to be artificially high due to the inclusion of uninteresting cases containing the "too short" keyword.

As mentioned above, we are interested in the identification of three types of cholesterol tests. To achieve this objective, in what follows we focus on a set of four keywords (totale, colesterolo, ldl, hdl) with three identifying letters. We also specify the newcode() option to generate a new variable recoding the observations that match the specified keywords.

At this point, we describe more deeply the recoding mechanism of screening:

- If newcode() is specified, a new variable is generated, taking as values the position of the keywords or regular expressions specified through the keys() option. The coding process is driven by the order of keywords or regular expressions.
- If recode() is specified, the newcode() newvar suboption is recoded according to the user-defined coding scheme.

Thus a first recoding of the source variable can be obtained as follows:

- . screening, sources(diagn\_test\_descr, lower)
- > keys("totale" "colesterolo" "ldl" "hdl") letters(3 3 3 3) explore(count)
- > newcode(tmp\_diagn\_test\_code)

Source	Key	Freq.	Percent
diagn_test_descr	tot	7304	29.47
	col	12595	50.81
	ldl	2015	8.13
	hdl	2872	11.59
	Total	24786	100.00

. tabulate tmp\_diagn\_test\_code

tn	np_diagn_t est_code	Freq.	Percent	Cum.
	1 2 3 4	7,304 7,535 12 11	49.15 50.70 0.08 0.07	49.15 99.85 99.93 100.00
	Total	14,862	100.00	

The explore(count) option instructs screening to display a table of frequency counts of all matched cases. The newcode() option creates tmp\_diagn\_test\_code, which is a new variable that takes as values the position of the keywords or regular expressions specified through the keys() option. The coding process is driven by the order of keywords or regular expressions: the number 1 is associated with the 7,304 observations matching the first keyword, tot; the number 2 is associated with the 7,535 observations matching the second keyword, col; and so on. Hence, by specifying keys("totale" "colesterolo" "ldl" "hdl") together with letters(3 3 3 3), tot takes precedence over col in the recoding process. This means that if some observations are recoded according to the first keyword match, they will not be recoded according to the following keywords in the keys() list, even if they match.

For this reason, the best recoding strategy is to first specify keywords that uniquely identify the cases of interest. Because keywords hdl and ldl each uniquely identify a cholesterol test, they must have priority in the recoding process over totale, which is an extension common to other pathologies.

Indeed, when we reverse the order of the keywords and specify the replace suboption in the newcode() option, screening produces

- . screening, sources(diagn\_test\_descr, lower)
- > keys("hdl" "ldl" "colesterolo" "totale") letters(3 3 3 3)
- > newcode(tmp\_diagn\_test\_code, replace)

WARNING: By specifying -replace- sub-option you are overwriting the -newcode()-> variable.

. tabulate tmp\_diagn\_test\_code

tmp_diagn_t est_code	Freq.	Percent	Cum.
1 2 3	2,872 2,015 7,731	19.32 13.56 52.02	19.32 32.88 84.90
4	2,244	15.10	100.00
Total	14,862	100.00	

where the newcode() variable now identifies all hdl and ldl cases. Notice that here we followed the correct approach, from specific to general. Moreover, as shown by the following code, when we specify the newcode() suboption label, screening attaches the specified keywords as value labels to the newcode() variable.

- . screening, sources(diagn\_test\_descr, lower)
- > keys("hdl" "ldl" "colesterolo" "totale") letters(3 3 3 3)
- > newcode(tmp\_diagn\_test\_code, replace label)

WARNING: By specifying -replace- sub-option you are overwriting the -newcode()-> variable.

. tabulate tmp\_diagn\_test\_code

tmp_diagn_t est_code	Freq.	Percent	Cum.
hdl	2,872	19.32	19.32
ldl	2,015	13.56	32.88
colesterolo	7,731	52.02	84.90
totale	2,244	15.10	100.00
Total	14,862	100.00	

The last step toward recoding is achieved by using the recode() option. This option allows you to recode the newcode() variable according to a user-defined coding scheme. When you specify this option, the coding process is completely under your control. The recode() option requires a recoding\_rule followed by a "user\_defined\_code" (the "user\_defined\_code" must be enclosed within double quotes).

When we specify recode(1 "90.14.1" ...), the standard code "90.14.1" will be used to recode all matched cases from the first keyword (hdl); when we specify

recode(... 2 "90.14.2" ...), the standard code "90.14.2" will be used to recode all matched cases from the second keyword (ldl); and so on. The third and forth keywords deserve special attention. totale (which was specified as the forth keyword, hence position 4) is a common extension that we want to identify only when it is matched simultaneously with colesterolo (which was specified as the third keyword, hence position 3). Thus the appropriate syntax in this case will be recode(... 3,4 "90.14.3" ...). Finally, when we specify recode(... 3 "not class. tests"), the code "not class. tests" will be used to recode all matched cases from the third keyword (colesterolo) that are not classified because they do not contain any further specification.

The final syntax of our example is

- . screening, sources(diagn\_test\_descr, lower)
- > keys("hdl" "ldl" "colesterolo" "totale") letters(3 3 3 3)
- > newcode(diagn\_test\_code)
- > recode(1 "90.14.1" 2 "90.14.2" 3,4 "90.14.3" 3 "not class. tests")
- . tabulate diagn\_test\_code

diagn_test_code	Freq.	Percent	Cum.
90.14.1	2,872	22.76	22.76
90.14.2	2,015	15.97	38.73
90.14.3	5,055	40.06	78.79
not class. tests	2,676	21.21	100.00
Total	12,618	100.00	

As the tabulate command shows, the new variable diagn\_test\_code is created according to the user-defined codes. Notice that only 5,055 cases are coded as "total cholesterol" (90.14.3). A two-way tabulate command (below) helps to highlight that 2,244 cases have to be considered incorrect identifications—that is, cases containing the same spelling of the keywords (totale) but related to other types of diagnostic tests<sup>4</sup>—whereas 2,676 are incomplete because they contain only colesterolo without further specification.

. tabulate diagn\_test\_code tmp\_diagn\_test\_code if tmp\_diagn\_test\_code !=., m

		tmp_diagn	_test_code		
diagn_test_code	hdl	1d1	colestero	totale	Total
	0	0	0	2,244	2,244
90.14.1	2,872	0	0	0	2,872
90.14.2	0	2,015	0	0	2,015
90.14.3	0	0	5,055	0	5,055
not class. tests	0	0	2,676	0	2,676
Total	2,872	2,015	7,731	2,244	14,862

This example shows that **screening** is a simple tool to manage complex string variables. Once you have obtained structured data (in our example, a categorical variable indicating cholesterol tests), you can finally start your statistical analysis.

<sup>4.</sup> Because of space restrictions, we deliberately omit the tabulation of such cases. It is available upon request.

#### 5 Extensions

Although the main utility of screening is the direct translation of complex narrative-text variables in a user-defined coding scheme, the command is flexible enough to cover a wide range of situations. In section 5.1, we present an example of how to use the command to facilitate the merging of information from different sources, while in section 5.2, we show how to use screening to extract or rearrange a portion of a string variable.

#### 5.1 Merging from different sources

In applied studies, a classic problem comes from trying to merge information from different sources that use different codes for the same units. A recently released command, kountry (Raciborski 2008), is an important step toward a solution.

The kountry command can be used to facilitate the merging of information from different sources by recoding a string variable into a standardized form. This recoding is possible using a custom dictionary created through a helper command.<sup>5</sup> In this section, we show an alternative way to merge information from different sources by using the screening command.

As an example, we try to merge two Italian datasets, one provided by the National Statistical Office (National Institute of Statistics in Italy) and the other provided by the Italian Ministry of the Interior. The two datasets contain, for each Italian municipality, the complete name and an alphanumeric code, the latter being different across sources. In theory, with the (uniquely identified) name of each municipality, it should be easy to merge the two datasets.

We first proceed by matching the two original datasets:

- . use istat, clear
- . sort comune
- . merge m:m comune using ministero
   (output omitted)
- . tabulate \_merge

_merge	Freq.	Percent	Cum.
master only (1) using only (2) matched (3)	288 290 7,812	3.43 3.46 93.11	3.43 6.89 100.00
Total	8,390	100.00	

<sup>5.</sup> See help kountryadd (if kountry is installed).

As you can see, there are 288 inconsistencies.<sup>6</sup> When we tabulate the unmatched cases, we would realize that unconventional expressions, like apostrophes, accents, double names, etc., are responsible for this imperfect result:

- . preserve
- . sort comune
- . drop if \_merge==3
  (7812 observations deleted)
- . list comune \_merge in 1/20, separator(20) noobs

comune	_merge
AGLIE'	2
AGLI	1
ALA´ DEI SARDI	2
ALBISOLA MARINA	2
ALBISOLA SUPERIORE	2
ALBISSOLA MARINA	1
ALBISSOLA SUPERIORE	1
ALI	2
ALI TERME	2
ALLUVIONI CAMBIO'	2
ALLUVIONI CAMBI	1
ALME´	2
ALM	1
AL DEI SARDI	1
AL	1
AL TERME	1
ANTEY-SAINT-ANDRE	2
ANTEY-SAINT-ANDR	1
APPIANO SULLA STRADA DEL	2
APPIANO SULLA STRADA DEL VINO	1

. restore

If you wish to recover all 288 unmatched municipalities, the proposed command is a simple and fast solution. Indeed, when you take advantage of the available options, you can (almost) completely recover unmatched cases with only one command. As an example, we recover nine cases (it is possible to recover all cases with this procedure), with a loop running on values of <u>merge</u> equal to 1 or 2, that is, running only on unmatched cases:

<sup>6.</sup> The number of unmatched cases is different between the master (288) and the using (290) datasets because of aggregation and separation of municipalities. Solving this kind of problem is beyond the illustrative scope of this example.

```
. forvalues i=1/2 {
 2.
            preserve
            keep if _merge==`i´
 3.
 4.
. screening, sources(comune) keys("ALBISSOLA" "AQUILA D'ARROSCIA" "BAJARDO"
> "BARCELLONA" "BARZAN" "BRIGNANO" "CADERZONE" "CAVAGLI" "MARINA" "SUPERIORE")
> cases(cases) newcode(comune, replace)
> recode(1,9 "ALBISOLA MARINA" 1,10 "ALBISOLA SUPERIORE" 2 "AQUILA DI ARROSCIA"
> 3 "BAIARDO" 4 "BARCELLONA POZZO DI GOTTO" 5 "BARZANO" 6 "BRIGNANO FRASCATA"
> 7 "CAVAGLIA" 8 "CADERZONE TERME")
            if `i'==1 drop codice_ente
 5.
            if `i'==2 drop codice
 7.
            keep comune codice
           sort comune
 8.
            save new_`i´,replace
 9.
 10.
           restore
11. }
(8102 observations deleted)
WARNING: By specifying -replace- sub-option you are overwriting the -newcode()-
> variable.
(note: file new_1.dta not found)
file new_1.dta saved
(8100 observations deleted)
WARNING: By specifying -replace- sub-option you are overwriting the -newcode()-
(note: file new_2.dta not found)
file new 2.dta saved
. keep if _merge==3
(578 observations deleted)
. save perfect_match, replace
(note: file perfect_match.dta not found)
file perfect_match.dta saved
. use new_1, clear
. merge 1:1 comune using new_2
 (output omitted)
. tabulate _merge
```

_merge	Freq.	Percent	Cum.
master only (1) using only (2) matched (3)	279 281 9	49.03 49.38 1.58	49.03 98.42 100.00
Total	569	100.00	

- . append using perfect\_match
- . tabulate \_merge

_merge	Freq.	Percent	Cum.
master only (1) using only (2)	279 281	3.33 3.35	3.33 6.68
matched (3)	7,821	93.32	100.00
Total	8,381	100.00	

Because we deliberately recovered only nine cases, the number of unmatched cases before the execution of screening is improved by nine cases, from 7,812 to 7,821 exact matches.

#### 5.2 Extracting a piece of a string variable

In this section, we show through three examples how screening can be used to extract or rearrange a portion of a string variable.<sup>7</sup>

#### Example 1

Imagine you have the string variable address, and you want to create a new variable that contains just the zip codes. Here is what the source variable address may look like:

. list, noobs sep(10)

```
address

4905 Lakeway Drive, College Station, Texas 77845 USA
673 Jasmine Street, Los Angeles, CA 90024
2376 First street, San Diego, CA 90126
66666 West Central St, Tempe AZ 80068
12345 Main St. Cambridge, MA 01238-1234
12345 Main St Sommerville MA 01239-2345
12345 Main St Watertwon MA 01239 USA
```

To find the zip code, you have to use screening with specific regular expressions, allowing it to exactly match all cases in the source variable address. Some examples of specific regular expressions are the following:

- ([0-9][0-9][0-9][0-9]) to find a five-digit number, the zip code
- [\-]\* to match zero or more dashes, or --
- [0-9]\* to match zero or more numbers, that is, the zip code plus any other numbers
- [ a-zA-Z]\* to match zero or more blank spaces and (lowercase or uppercase) letters

Once the correct regular expression(s) is found, to use screening to create a new variable containing the zip codes, you have to do the following:

<sup>7.</sup> The following examples have been taken from the UCLA website resources to help you learn and use Stata. See http://www.ats.ucla.edu/stat/stata/faq/regex.htm.

- 1. Use the newcode() option to create the new variable zipcode.
- 2. Combine the above regular expressions and use them as a unique keyword.
- 3. Use the regexs(n) function as a "user\_defined\_code" in the recode() option. regexs(n) returns the subexpression n from the respective keyword match, where  $0 \le n \le 10$ . Stata regular-expression syntaxes use parentheses, (), to denote a subexpression group. In particular, n = 0 is reserved for the entire string that satisfied the regular expression (keyword); n = 1 is reserved for the first subexpression that satisfied the regular expression (keyword); and so on.

#### Hence, you may code

```
. screening, sources(address)
> keys("([0-9][0-9][0-9][0-9])[\-]*[0-9]*[ a-zA-Z]*$")
> cases(c) newcode(zipcode) recode(1 "regexs(1)")
WARNING! You are SCREENING some keywords using regular-expression operators
> like ^ . ( ) [ ] ? *
```

- 1) Option -letter- doesn't work IF a keyword contains regular-expression operators
- 2) Unless you are looking for a specific regular-expression, regular-expression operators must be preceded by a backslash \ to ensure keyword-matching (e.g. \^\.)
- 3) To match a keyword containing \$ or \, you have to specify them as [\\$] [\\]
- . tabulate zipcode

Notice that:

zipcode	Freq.	Percent	Cum.
01238	1	14.29	14.29
01239	2	28.57	42.86
77845	1	14.29	57.14
80068	1	14.29	71.43
90024	1	14.29	85.71
90126	1	14.29	100.00
Total	7	100.00	

#### where recode(1 "regexs(1)") indicates that

- 1. 1 is the *recoding\_rule*; that is, the coding process is related to the first (and unique) keyword match.
- 2. regexs(1) is used to recode. Indeed, it returns the string related to the first (and unique) subexpression match.<sup>8</sup>

As a result, the new variable **zipcode** is created by using only one line of code. Notice that **screening** warns you that you are matching a keyword containing one or more regular-expression operators.

<sup>8.</sup> Remember that subexpressions are denoted by using (). In the considered syntax, the only subexpression is represented by ([0-9][0-9][0-9][0-9]). This means that, in this case, you cannot specify n > 1.

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#### Example 2

Suppose you have a variable containing a person's full name. Here is what the variable fullname looks like:

. list, noobs sep(10)



Our goal is to swap first name with last name, separating them by a comma. The regular expression to reach the target is (([a-zA-Z]+)[]\*([a-zA-Z]+)). It is composed of three parts:

- 1. ([a-zA-Z]+) to capture a string consisting of letters (lowercase and uppercase), that is, the first name
- 2. []\* to match with a space(s), that is, the blank between first and last name
- 3. ([a-zA-Z]+) again to capture a string consisting of letters, this time the last name

The following is a way to proceed using screening:

```
fullname

Adams, John
Smiths, Adam
Smiths, Mary
Wade, Charlie
```

Notice the newcode() suboption add. It can be specified only when a regexs(n) function is specified as a "user\_defined\_code" in the recode() option. The add suboption allows for the creation of the newcode() variable as a concatenation of subexpressions returned by regexs(n). In the example above,

- 1. recode(1 "regexs(2)," ... returns the second subexpression from the first keyword match (the last name) plus a comma.
- 2. ...2 "regexs(0)" ... returns the blank matched by the second keyword;
- 3. ...3 "regexs(1)") returns the first subexpression from the third keyword match (the first name).

As a result, the variable fullname is replaced (note the suboption replace) sequentially by the concatenation of subexpressions returned by 1, 2, and 3 above.

#### Example 3

Imagine that you have the string variable date containing dates:

. list date, noobs sep(20)



The goal is to produce a string variable with the appropriate four-digit year for each case, which Stata can easily convert into a date. You can achieve the target by coding something like the following:

```
. generate day = regexs(0) if regexm(date, "^[0-9]+")
. generate month = regexs(0) if regexm(date, "[a-zA-Z]+")
. generate year = regexs(0) if regexm(date, "[0-9]*$")
. replace year = "20"+regexs(0) if regexm(year, "^[0][0-9]$")
(2 real changes made)
. replace year = "19"+regexs(0) if regexm(year, "^[1-9][0-9]$")
(2 real changes made)
. generate date1 = day+month+year
```

list.	noobs	sep(	10)

month	year	date1
jan	2007	20jan2007
June	2006	16June2006
sept	1985	06sept1985
june	2004	21june2004
july	1990	4july1990
jan	1999	9jan1999
aug	1999	6aug1999
august	2003	19august2003
	jan June sept june july jan aug	jan 2007 June 2006 sept 1985 june 2004 july 1990 jan 1999 aug 1999

Alternately, you can obtain the same result by using screening:

```
. screening, sources(date) keys("^[0-9]+" "[a-zA-Z]+" "[0][0-9]$" "[1-9][0-9]$")
> newcode(date1, add)
> recode(1 "regexs(0)" 2 "regexs(0)" 3 "20+regexs(0)" 4 "19+regexs(0)")
WARNING! You are SCREENING some keywords using regular-expression operators
> like ^ . ( ) [ ] ? *
    Notice that:
```

- 1) Option -letter- doesn't work IF a keyword contains regular-expression operators
- 2) Unless you are looking for a specific regular-expression, regular-expression operators must be preceded by a backslash \ to ensure keyword-matching (e.g. \^\.)
- 3) To match a keyword containing \$ or \, you have to specify them as [\\$] [\\]
- . list date date1, noobs sep(10)

date	date1
20jan2007	20jan2007
16June06 06sept1985	16June2006 06sept1985
21june04	21 june 2004
4july90	4july1990
9jan1999	9jan1999
6aug99	6aug1999
19august2003	19august2003

Also in this case, as in the previous example, we specify the newcode() suboption add because we need to create the newcode() variable as a concatenation of subexpressions from keyword matching. The same result can be obtained using the following syntax:

(Continued on next page)

- 1) Option -letter- doesn't work IF a keyword contains regular-expression operators
- 2) Unless you are looking for a specific regular-expression, regular-expression operators must be preceded by a backslash \ to ensure keyword-matching (e.g. \^\\.)
- 3) To match a keyword containing \$ or \, you have to specify them as [\\$] [\\]
- . list date date1, noobs sep(10)

date	date1
20jan2007 16June06	20jan2007 16June2006
06sept1985	06sept1985
21june04 4july90	21june2004 4july1990
9jan1999	9jan1999
6aug99 19august2003	6aug1999 19august2003

where the only difference is represented by the way in which the *matching\_rule* is specified: begin instead of ^ and end instead of \$.

# 6 Summary

In this article, we introduced the new screening command, a data-management tool that helps you examine and treat the content of string variables containing free, possibly complex, narrative text. screening allows you to build new variables, to recode new or existing variables, and to build a set of categorical variables indicating keyword occurrences (a first step toward textual analysis). Considerable efforts were devoted to making the command as flexible as possible; thus screening contains a rich set of options that is intended to cover the most frequently encountered problems and necessities. Because of this flexibility, the command can be used in many different fields, like EPR data, data from different sources, or survey data. The execution of screening is fast, thanks to Mata programming; its syntax is simple and common to many other Stata commands, thus it is useful for all users regardless of their levels of experience in Stata. We especially recommend that you use the explore() option; it makes the command a useful data-mining tool. Nevertheless, expert users can exploit a more complicated syntax that substantially eases the preparatory burden for data cleaning.

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#### About the authors

Federico Belotti is a PhD student in econometrics and empirical economics at the University of Rome Tor Vergata.

Domenico Depalo is a researcher in the Economic Research Department of the Bank of Italy in Rome. He received his PhD in econometrics and empirical economics from the University of Rome Tor Vergata and was enrolled in a Post Doc program at the University of Rome La Sapienza.