BMJ Open Health literacy in Italy: a cross-sectional study protocol to assess the health literacy level in a population-based sample, and to validate health literacy measures in the Italian language

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ABSTRACT

Introduction Health literacy (HL) concerns the knowledge and competences necessary for people to meet complex health demands. The aims of this study are to assess the level of HL in a sample using the Italian version of the Newest Vital Sign and the association of HL and selected antecedents with health outcomes, and to develop and validate the Italian version of the three Brief Health Literacy Screeners, two subjective numeracy items and the short form and the short-short form of the European Health Literacy Survey Questionnaire.

Methods and analysis The study adopts a cross-sectional design and is being conducted in Florence, with information collected through telephone interviews. The population-based sample has been randomly selected using the registries of eight general practitioners (GPs). Based on a power calculation, 480 subjects will be included. Participants have been randomly offered two different questionnaires, each containing different HL measures. Data on sociodemographics and important antecedents and consequences of HL will be collected and the distribution of HL levels calculated. The mediating role of HL will be assessed using Preacher and Hayes' model. To assess the concurrent validity of the HL scales, correlation and receiver operating characteristic analyses will be performed.

Ethics and dissemination The study protocol has been approved by the Ethics Committee of the Area Vasta Centro. Results will be disseminated via scientific journals and conference presentations, and individual data made available to the GPs.

BACKGROUND

Health literacy (HL) concerns the knowledge and competences of individuals necessary for meeting the complex health demands of modern society. According to the conceptual framework of Sørensen *et al* 'HL is linked to literacy and entails people's knowledge, motivation and competences to access, understand, appraise and apply health information

Strengths and limitations of this study

- This is the first study in Italy to attempt to assess health literacy and its related antecedents and outcomes, and to validate a variety of different measures.
- The use of numerous measures will enable the systematic comparison of data with those obtained in other countries.
- The direct involvement of general practitioners (GPs) who signed the information sheet and agreed with the aims of the study could lead to high compliance.
- One limitation of this study is that the sample has not been randomly selected among people living in Florence, but from individuals registered as patients with different GPs.
- Another limitation is that the GPs have been recruited using convenience criteria, potentially introducing a selection bias.

in order to make judgements and take decisions in everyday life concerning healthcare, disease prevention and health promotion to maintain or improve quality of life during the life course'.¹ In their framework, the authors identified 12 dimensions of HL, as well as proximal and distal factors (antecedents) that influence HL and related outcomes (consequences). Distal factors include societal and environmental factors, whereas proximal factors are situational and personal determinants such as general literacy, individual characteristics, and prior experience with illness and the healthcare system. On the other hand, the consequences at the individual and population levels refer to health service use and health costs, health behaviour and health outcomes, participation and empowerment in health issues, and equity and sustainability

of public health issues. Despite this and other conceptual models (see, eg, the models of Nutbeam,² Paasche-Orlow and Wolf,³ and Schulz and Nakamoto⁴), to date the causal relationships between antecedents, HL and health outcomes are still unclear. Only a few studies have systematically investigated the role of HL as a mediator between antecedents and different health outcomes, while many authors have described HL as a cofactor or confounder in predicting health outcomes.^{5–13}

Many tools exist to measure HL, but to date none of them are considered the gold standard.^{14–15} Measures largely vary in their approach and design, as well as in terms of their purpose. Some of them are screening tools that were originally developed for the clinical setting—to quickly identify patients with lower HL levels. Other tools were developed for the purpose of measuring the broader concept of HL to provide an in-depth assessment of the different dimensions of HL and exploring its relationship with determinants or outcomes. Moreover, HL measures are either performance based (objective) or self-reported (subjective).^{14–17} This means that they evaluate an individual's HL level by asking individuals to either directly apply their skills to a healthcare-related task or report on their skills, such as being confident enough to fill out medical forms on their own (see, eg, the studies of Chew *et al*^{18–19}). Overall, there has been relatively little systematic research comparing the effectiveness of performance-based and self-reported measures, specifically with regard to possible relationships between HL and health outcomes. As suggested in a recent review, further studies that assess HL by using performance-based and self-reported tools at the same time are needed to clarify these aspects.²⁰

In Italy, research on HL is still relatively new; only a few studies have assessed and published results on HL in Italian samples.^{21–22} Therefore, the aims of this study are:

1. to assess the level of HL in a population-based sample in Florence, Italy, using the Italian version of the Newest Vital Sign (NVS-IT);

2. to assess the association of HL, selected antecedents and health outcomes, so as to evaluate whether HL is a mediator, a confounding factor or a cofactor in predicting the outcome variables;
3. to develop and validate the Italian version of three subjective Brief Health Literacy Screeners (BHLS) and two subjective numeracy items, and to assess their diagnostic accuracy in detecting subjects with limited HL in comparison to the NVS-IT (concurrent validation);
4. to develop and validate the Italian version of the short form (HLS-EU-Q16) and short-short form (HLS-EU-Q6) of the European Health Literacy Survey Questionnaire (HLS-EU), and to assess their diagnostic accuracy in detecting subjects with limited HL in comparison to the NVS-IT (concurrent validation).

METHODS

The study adopts a cross-sectional design and is being conducted in Florence, Italy and its surroundings. It started in February 2017 and will presumably end in November 2017, with information collected through telephonic interviews.

Study population and sampling criteria

The population-based sample was randomly selected from the registries of eight general practitioners (GPs) working in primary healthcare centres of the municipality of Florence—a sampling method that is also suggested by other authors (see, eg, the study of Toçi *et al*²³) (figure 1). According to the regulations of the National Healthcare System and the Constitution of the Italian Republic, in Italy every citizen and foreign resident over the age of 18 has the right to be registered as a patient with a GP. Healthcare is provided by GPs free of charge to all patients. In the municipality of Florence, about 200

Figure 1 Sampling procedure. GP, general practitioner.

GPs provide continuing and comprehensive healthcare to individuals who are registered as patients. For each GP, the number of registered and assisted patients is on average 1100, with a maximum limit of 1500 patients per doctor.

The GPs were recruited using convenience criteria. Both the President of the Provincial Medical Council and the representative of the GPs at the University Hospital of Florence informed their colleagues to join the study via either email or verbal contact. The first eight, to voluntarily join the study, were included. Each GP randomly selected 80 subjects among those registered as one of his/her patients.

Inclusion criteria were the following: 18–69 years of age and Italian speaking (since the survey is conducted in Italian). Exclusion criteria included cognitive impairment, severe psychiatric diseases and end-stage diseases. Each GP verified the inclusion and exclusion criteria when selecting the sample.

Sample size was calculated based on the first aim of this study. Since no data on HL in Italy using Newest Vital Sign (NVS) as a measure is publicly currently available, data from the HLS-EU were used as a reference. Based on the original study that assessed HL across eight different European countries (Italy not included),⁶ Spain—with the highest rate of limited HL when assessed with the NVS—was chosen (34.3%) as the expected value. Considering a CI of 95% and a margin of error equal to 0.05, the sample size was established as 480 subjects. An oversampling of 160 individuals is planned to account for problems in recruitment (ie, subjects not available for telephonic interviews or refusal rates). If oversampling is not sufficient, a second random sample will be drawn.

Each subject is randomly allocated to one of the two arms of the study (A and B), according to the questionnaires used during the interview (type I and type II questionnaires, respectively).

Procedures: data collection

Each selected subject is contacted via postal mail. Subjects receive an information sheet signed by both the GP and the person in charge of the study, which includes a short description of the study, an invitation to participate and a consent form. Participants are asked to sign the consent form and return it via mail to the researchers in charge. The mail also contains the nutritional label of the NVS-IT. For further information, email addresses and telephone numbers of the researchers are provided in the letter. After receipt of the signed consent forms, the subjects are contacted over phone for the interview.

Nine interviewers who are part of the research group make the phone calls. Written instructions on how to conduct the interview are drawn up and shared within the research team so as to standardise the procedure and limit interviewer bias. Each subject is randomly assigned to one of the nine interviewers and contacted a maximum of six times before being considered unreachable.

Table 1 Summary of the content of the two types of questionnaires that will be used in the study

Questionnaire sections	Questionnaire type I	Questionnaire type II
General section	X	X
Health literacy measurements		
NVS-IT	X	X
The three-item BHLS	X	
Two subjective numeracy items	X	
HLS-EU-Q16 and HLS-EU-Q6		X
Health outcome variables	X	X

BHLS, Brief Health Literacy Screener; HLS-EU-Q6, short-short form of the European Health Literacy Survey Questionnaire; HLS-EU-Q16, short form of the European Health Literacy Survey Questionnaire; NVS-IT, Italian version of the Newest Vital Sign.

Data are collected using an electronic database and anonymised by assigning a numeric code to each recruited person.

To meet the aims of the study, two questionnaires (type I and type II) have been drawn up, each containing the sections on the possible antecedents and consequences of HL, as well as different items to assess HL (table 1). Both questionnaires have a general section that includes questions on sociodemographics, familial data (antecedents) and health-related outcomes. In addition, both questionnaires include the NVS-IT items. However, whereas the type I questionnaire has three BHLS and two subjective numeracy items, the type I questionnaire contains a section with the HLS-EU-Q16.

The questionnaires have both been pretested with a small sample of volunteers, and take about 20–25 min, depending on the questionnaire.

Measurements

Newest Vital Sign

The NVS is a commonly used objective measure of HL. It was originally developed in the USA for English speakers and Spanish speakers.²⁴ However, in recent years, the NVS has seen increased application in countries other than the USA,^{6 25–30} including Italy (NVS-IT).³¹ Moreover, it was originally developed to be administered by face-to-face interviews, although in one study, a web-based survey was conducted²⁹ and in another it was self-administered.³² To the best of our knowledge, no published studies have reported NVS data collected through telephonic interviews.

The NVS-IT consists of an ice cream nutrition label, with seven associated questions that measure literacy and numeracy. It produces a final score ranging from 0 to 6, allowing subjects to be classified in three categories—high likelihood of limited HL (score: 0–1), possibility of limited HL (score: 2–3) and adequate HL (score: 4–6). This instrument takes very little time to be administered

(3–5 min), and has been found to be easily applicable to people in different settings.^{33 34} Since the information needed to answer the questions has to be derived from the nutritional label, the label is included in the postal mail to be sent to the GP-assisted participants.

HL and numeracy screeners

In addition to the NVS-IT, the type I questionnaire also contains items of two other measurement tools—three BHLS and two subjective numeracy items. Both measures are self-reported with Likert-type responses, validated for predicting NVS scores by a previous study.³⁵

The three BHLS were initially developed and validated in English.¹⁸ They assess literacy, interaction, comprehension and confidence (self-efficacy) skills. Even though they were originally developed for the clinical setting, they have seen increased application in population-based studies.^{36–41}

The two subjective numeracy items—adapted from the STAT-confidence that assesses people's confidence in understanding medical statistics⁴²—were initially also validated in English. The two questions assess an individual's confidence in medical statistics by evaluating numeracy, comprehension, application/function, decision-making and confidence. Both tools produce a score to measure the HL levels.

HLS-EU-Q16 and HLS-EU-Q6

In addition to the NVS-IT, the type II questionnaire contains the HLS-EU-Q16 and the HLS-EU-Q6 which are respectively the short form and the short-short form of the 47-item European Health Literacy Survey Questionnaire (HLS-EU-Q47). The HLS-EU was developed in the aim of measuring and comparing HL levels of populations in different European countries. Based on the conceptual model proposed by Sørensen *et al*,¹ a European Union-funded working group developed the HLS-EU-Q47.⁴³ It includes 47 items covering 12 subdomains (including domains such as accessing and obtaining, understanding or appraising information relevant to healthcare, disease prevention and health promotion). It is a self-reported tool with Likert-type responses ('very easy', 'fairly easy', 'fairly difficult', 'very difficult') and an associated final score that measures interaction, comprehension, information seeking, application/function, decision-making/critical thinking, evaluation, responsibility, confidence and navigation skills. Based on the extended version, two short versions were developed by selecting 16 items for the HLS-EU-Q16 and 6 items for the HLS-EU-Q6. Even though the reduction in the number of items has decreased the informative value and theoretical scope of the measure, it has increased its utility by reducing its administration time, while maintaining strong correlations to the full instrument.⁴⁴

To generate the score for the HLS-EU-Q16, answers are dichotomised ('don't know' answers are coded as missing values). 'Fairly difficult' and 'very difficult' are both coded with 0 (zero), whereas 'fairly easy' and 'very

easy' are both coded with 1. The HLS-EU-Q16 score is a sum score (range: 0–16) and according to the final score, three levels of HL have been defined: inadequate HL (0–8); problematic HL (9–12); sufficient HL (13–16).

Scores for the HLS-EU-Q6 are calculated similarly to those of the HLS-EU-Q47 (from 1 for 'very difficult', to 4 for 'very easy'). The final score is the mean score that is calculated if at least five of the six items are completed and range from 1 to 4. According to the final score, three possible levels of HL have been defined: inadequate HL (1–2); problematic HL (2–3); sufficient HL (3–4).

Translation of HL measurements

The English versions of the BHLS, two subjective numeracy items, the HLS-EU-Q16 and the HLS-EU-Q6 have been translated and adapted using standard procedures, including forward and backward translation (performed by Italian and English native speakers). A final Italian version was drafted and then shared and discussed with members of the research group.⁴⁵ Table 2 reports on the Italian versions of the tools.

Antecedents

Data on antecedents of HL were collected according to the proposal of the HLS-EU⁶ and the Italian lifestyle surveillance system PASSI (Progress by Local Health Units towards a Healthier Italy).⁴⁶ In addition to including questions on gender, birth year, nationality, number of years living in Italy (for those who were born abroad), educational level and marital status, the questionnaire also has questions on the number of family members living in the same household, whether one has ever received training or is/has been employed in the field of healthcare, employment status (currently having a paid job), financial situation, and whether a family member or a friend normally accompanies him/her to medical appointments.

Health outcome variables

According to the Sørensen model¹, many outcomes are associated with HL. In line with what has been already made public by the HLS-EU,⁶ the following information will be collected: self-assessed health status (excellent, very good, good, so-so/fair, bad), weight and height to calculate body mass index, health services used in the last 12 months (number of doctor visits, hospital admissions, emergency department visits and use of healthcare services such as dentists, physiotherapists, psychologists, dieticians or opticians. Responses are coded as follows: 0, 1–2 times, 3–5 times, 6 times or more, don't know/refusal). To measure health status, subjects are asked about long-term illnesses (illnesses that have lasted or are expected to last for at least 6 months), coded in four categories: yes, more than one; yes, one; no; don't know/refusal.

Statistical analysis

Statistical analysis will be performed using IBM SPSS Statistics for Windows, V.24.0 (IBM) and StataIC V.11

Table 2 The three-item HL screening, the two subjective numeracy items, the HLS-EU-Q16 and the HLS-EU-Q6 with answers and scoring (English and Italian versions)

Measurement tool	English version	Italian version
Three-item BHLS	<p>1. How often do you have problems learning about your medical condition because of difficulty in understanding written information? Always (1); often (2); sometimes (3); occasionally (4); never (5)</p> <p>2. How often do you have someone help you read hospital materials? Always (1); often (2); sometimes (3); occasionally (4); never (5)</p> <p>3. How confident are you in filling out medical forms by yourself? Extremely (1); quite a bit (2); somewhat (3); a little bit (4); not at all (5)</p>	<p>1. Quanto spesso ha difficoltà a comprendere informazioni scritte che riguardano il suo stato di salute? Sempre (1); spesso (2); a volte (3); raramente (4); mai (5)</p> <p>2. Quanto spesso ha qualcuno (un familiare, un amico o un'altra persona) che la aiuta a leggere il materiale sanitario? Sempre (1); spesso (2); a volte (3); raramente (4); mai (5)</p> <p>3. Quanto si sente sicuro nel compilare da solo moduli medici? Moltissimo (1); Molto (2); abbastanza (3); poco (4); per niente (5)</p>
Two subjective numeracy items	<p>1. In general, how easy or hard do you find it to understand medical statistics? Very easy (1); easy (2); hard (3); very hard (4)</p> <p>2. How much do you agree or disagree with the following statement? In general, I depend on numbers and statistics to help me make decisions about my health. Strongly agree (1); somewhat agree (2); somewhat disagree (3); strongly disagree (4)</p>	<p>1. In generale, quanto è facile o difficile per lei comprendere risultati di indagini statistiche condotte in ambito medico? Molto facile (1); facile (2); difficile (3); molto difficile (4)</p> <p>2. Quanto è d'accordo o in disaccordo con la seguente affermazione? In generale, mi baso su numeri e statistiche per prendere decisioni sulla mia salute. Totalmente d'accordo (1); abbastanza d'accordo (2); abbastanza in disaccordo (3); molto in disaccordo (4)</p>
HLS-EU-Q16 and HLS-EU-Q6 (*)	<p>On a scale from very easy to very difficult, how easy would you say it is to...</p> <p>Very easy; fairly easy; fairly difficult; very difficult; (don't know)†</p> <p>1. ... find information on treatments of illnesses that concern you?</p> <p>2. ... find out where to get professional help when you are ill?</p> <p>3. ... understand what your doctor says to you?</p> <p>4. ... understand your doctor's or pharmacist's instruction on how to take a prescribed medicine?</p> <p>5. ... judge when you may need to get a second opinion from another doctor?*</p> <p>6. ... use information the doctor gives you to make decisions about your illness?*</p> <p>7. ... follow instructions from your doctor or pharmacist?</p> <p>8. ... find information on how to manage mental health problems like stress or depression?*</p> <p>9. ... understand health warnings about behaviour such as smoking, low physical activity and drinking too much?</p> <p>10. ... understand why you need health screenings?</p> <p>11. ... judge if the information on health risks in the media is reliable?*</p> <p>12. ... decide how you can protect yourself from illness based on information in the media?</p> <p>13. ... find out about activities that are good for your mental well-being?*</p> <p>14. ... understand advice on health from family members or friends?</p> <p>15. ... understand information in the media on how to get healthier?*</p> <p>16. ... judge which everyday behaviour is related to your health?</p>	<p>Su una scala da estremamente difficile a estremamente facile, secondo Lei, quanto è difficile...</p> <p>Molto difficile; abbastanza difficile; abbastanza facile; molto facile; (non so)†</p> <p>1. ...trovare informazioni sui trattamenti delle malattie che La riguardano?</p> <p>2. ...identificare dove trovare personale sanitario quando è malato/a?</p> <p>3. ...capire cosa Le dice il Suo medico?</p> <p>4. ...capire le istruzioni del Suo medico o del Suo farmacista su come assumere un farmaco che Le è stato prescritto?</p> <p>5. ...giudicare quando è necessaria una seconda opinione da parte di un altro medico?*</p> <p>6. ...usare le informazioni che il medico le fornisce per prendere delle decisioni a proposito della Sua malattia?*</p> <p>7. ...seguire le istruzioni del Suo medico o del Suo farmacista?</p> <p>8. ...trovare informazioni su come gestire problemi di salute mentale, come lo stress o la depressione?*</p> <p>9. ...capire avvertenze per la salute riguardanti comportamenti come fumare, fare poca attività fisica e bere troppo alcool?</p> <p>10. ...capire perché è necessario sottoporsi agli screening (come mammografia, PAP test, ricerca di sangue occulto nelle feci, determinazione glicemia, misurazione della pressione sanguigna)?</p> <p>11. ...giudicare se le informazioni sui rischi per la salute riferite dai mezzi di comunicazione siano attendibili?*</p> <p>12. ...decidere come può proteggersi da una malattia basandosi sulle informazioni riferite dai media?</p> <p>13. ...identificare attività che fanno bene al benessere mentale?*</p> <p>14. ...capire consigli da parte di familiari o amici riguardo alla salute?</p> <p>15. ...capire le informazioni riferite dai mezzi di comunicazione su come migliorare il suo stato di salute?*</p> <p>16. ...giudicare quali comportamenti quotidiani hanno un impatto sulla Sua salute?</p>

Continued

Table 2 Continued

Measurement tool	English version	Italian version
<p>†For HLS-EU-Q16, 'fairly difficult' and 'very difficult' are coded with 0 while 'fairly easy' and 'very easy' are coded with 1; for HLS-EU-Q6, the coding is as follows: Very easy=4; fairly easy=3; fairly difficult=2; very difficult=1; for HLS-EU-Q16 and HLS-EU-Q6, don't know is coded as 'missing'.</p> <p>BHLS, brief health literacy screeners; HL, health literacy; HLS-EU-Q6, short-short form of the European Health Literacy Survey Questionnaire; HLS-EU-Q16, short form of the European Health Literacy Survey Questionnaire.</p> <p>*Items included in the HLS-EU-Q6</p>		

(StataCorp). For each analysis, an alpha level of 0.05 will be considered as significant.

Prior to the main analysis, data will be assessed for non-normality.

Descriptive statistics will be calculated and data will be presented as percentage or mean $\pm$ SD. Associations between variables will be tested using Fisher's exact test for categorical data and the Student's t-test or analysis of variance for continuous data. Linear regression analysis will be performed to assess the linear relationships between HL and covariates. Receiver operating characteristic (ROC) analysis will be performed for validation. According to the aims of the study, different analyses will be carried out.

Aim 1. To assess the level of HL in a population-based sample in Florence, Italy, using the Italian version of the NVS-IT.

The distribution of HL levels will be calculated with the results projected back to the target population through inference analysis aimed at calculating estimates for the target population.

Aim 2. To assess the association of HL, selected antecedents and health outcomes, so as to evaluate whether HL is a mediator, a confounding factor or a cofactor in predicting the outcome variables.

Linear regression analyses will be performed to assess the linear relationship between NVS-IT scores, antecedents and health outcomes. In the first step, the association between antecedents (independent variables) and NVS-IT scores (dependent variable) will be assessed, followed by an assessment of the association between NVS-IT (independent variable) and health outcomes (dependent variable).

The potential mediating role of HL will be assessed using the Preacher and Hayes' model,⁴⁷ which is based on a normal theoretical approach and bootstrapping approach for obtaining reliable intervals between variables.^{48 49} This approach supports the traditional methods of Baron and Kenny⁴⁸ in studying the effects of mediating variables.⁵⁰

Aim 3. To develop and validate the Italian version of three BHLS and two subjective numeracy items, and to assess their diagnostic accuracy in detecting subjects with limited HL compared with the NVS-IT (concurrent validation).

Aim 4. To develop and validate the Italian version of the short form (HLS-EU-Q16) and short-short form (HLS-EU-Q6) of the HLS-EU, and to assess their diagnostic accuracy in detecting subjects with limited HL, compared with the NVS-IT (concurrent validation).

To assess the concurrent validity (the degree of agreement between two different tools in measuring the same concept) and the diagnostic accuracy of the HL scales, correlation (Pearson or Spearman) and ROC analyses will be performed. In order to verify that the thresholds suggested in the original validation studies are capable of distinguishing individuals with lower HL levels from individuals with higher HL levels, the sensitivity, specificity,

positive and negative predictive values and total accuracy will be calculated for each potential threshold value. Based on the already validated NVS-IT, two different analyses will be performed. The first analysis will consider participants with 'adequate HL' on the NVS-IT as being highly literate and those with 'possibility of limited HL' and 'high likelihood of limited HL' as being less literate. The second analysis will use a different grouping and assign those with 'Possibility of limited HL' on the NVS-IT to the highly literate group. The area under the ROC curve will be used as a measure of accuracy.⁵¹ To assess the level of agreement, Cohen's kappa will also be calculated.

Since the HLS-EU-Q16 has been developed from the HLS-EU-Q47 using Rasch analysis,⁴⁴ this analysis will be performed on the Italian version of the HLS-EU-Q16 to describe the 'test-item difficulty', its strengths and weaknesses.⁵²

Ethics and dissemination

The study has been designed following the principles of the Declaration of Helsinki. The protocol has been approved by the Ethics Committee of the 'Area Vasta Centro' (Local Health Unit of Central Tuscany, Careggi University Hospital and Meyer University Children's Hospital; Ref. CEAVC: 10113, 01 December 2016). As briefly described above, after data collection, information will be anonymised by assigning a numerical code to each participant. Any original identifiable information will be destroyed once the study has been completed.

Results will be disseminated through peer-reviewed scientific journals and conference presentations. Moreover, grouped data will be made available to GPs. Finally, the results will be discussed with policy-makers of the Tuscany Region.

DISCUSSION

Strengths and limitation

To the best of our knowledge, this is the first study in Italy that comprehensively attempts to assess HL and its related antecedents and outcomes and to validate a variety of different measures.

HL levels will be assessed in a population-based sample via use of a validated measure (NVS-IT), which has been applied in different countries and contexts^{25–30 32} and has also been used in the HLS-EU.⁶ This will allow for systematic comparison of data with those obtained in the eight countries involved in the European survey. Comparability will be made easier via use of a population-based sample of adults, similar to those included in the HLS-EU. In addition, given that the NVS-IT is already validated in Italian, the analysis will enable the systematic testing of the concurrent validity of the newly validated measures.

In this study, telephone interviews are used to collect NVS-IT data, while in almost all the studies face-to-face interviews are used. To date, no data have been published concerning the comparison of NVS data collected using different methods of administration. Compared with

face-to-face interviews, telephone interviews offer several advantages,⁴⁵ namely, the elimination of any bias caused by the appearance of the interviewer, lower costs related to the transfers of the interviewer or the person interviewed and lower administrative costs. Moreover, there is some evidence that people are more likely to report health-related events on the phone rather than in face-to-face interviews.⁴⁵ The potential problems with telephone interviewing are instead as follows: the respondent may seek help from another person at home, and there is no assurance who the person is at the other end on the line. This second risk is difficult to completely eliminate, while the first one is minor,⁴⁵ since the interviewer would be able to notice the involvement of other people (ie, the person on the phone would have to repeat each question or use the hands-free mode).

Another strength concerns the HL measures themselves, as they have already been validated in different languages and/or settings. Therefore, the diagnostic accuracy of the newly validated measures will be compared with that identified in other studies. In any case, as noted by Reeve and Basalik,⁵³ HL tests were primarily designed to detect illiteracy, so they often show a ceiling effect when used in the general population. The ceiling effect can significantly skew distribution, giving rise to concern about attenuated correlation. In our study, this phenomenon might be observed for NVS-IT, BHLS and two subjective numeracy items, while the HLS-EU-Q47 and its short forms have initially been validated in the general population. Any possible ceiling effect or skewed distribution will be considered in the statistical analysis.

Another strength of this study is the direct involvement of the GPs who have signed the information sheet and agreed with the aims of the study. Due to their strong involvement, participants might be more motivated to participate in the study, which is expected to result in higher compliance (see, eg, the study of Toçi *et al*²³).

One limitation of this study is that while the sample was randomly selected among people living in Florence, it includes individuals registered as patients with different GPs. Moreover, the GPs were recruited using convenience criteria, which might introduce a selection bias. To curb some of the bias, GPs were chosen according to their geographical location, in the aim of covering different districts in Florence that may present different characteristics of the residents. Furthermore, the variable that identifies the subject's GP, as well as the number of registered patients, will be considered and included as a confounder in the analyses.

Implications

This study has a number of important implications. First of all, the results of the study will be useful in assessing the ability of community members to seek, locate and use health services. Moreover, obtaining validated HL measures in Italian will support current research efforts on HL and related outcomes in Italy—both at a population-based level and in the clinical setting. As suggested

by other authors,⁴⁴ it is proposed to use the HLS-EU-Q6 for research other than HL-specific research in Italy. More specifically, the current plan is to integrate the HLS-EU-Q6 as a covariate in the Italian lifestyle surveillance system PASSI.

In addition, since the results will be presented and discussed with the Regional policy-makers, including the General Directors of Local and Hospital Health Units, data on HL will also be taken into account when planning community-based health and marketing activities, educational strategies to be used in educational campaigns or campaigns to support the introduction of new services, screening initiatives (eg, bowel or skin cancer) and vaccination programmes.⁵⁴

At a patient level, an important implication of the study relates to the direct involvement of the GPs. Apart from becoming more familiar with the concept of HL and its assessment tools, GPs will receive individual data for their patients, which will allow them to address findings in their day-to-day practice and potentially improve their care relationships.

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Patient consent Obtained.

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