PATTERNS OF USE OF ENGLISH SYNONYMOUS MEDICAL TERMS: THE CASE OF *DISEASE* AND *ILLNESS*. A CORPUS-BASED STUDY

by Barbara Cappuzzo^{*}

SUMMARY: 1. Abstract. - 2. Introduction.- 3. The case of *disease* and *illness.* - 3.1. Consulting generic bilingual dictionaries. - 3.2. Consulting specialized monolingual dictionaries. - 3.3. Consulting other dictionaries. - 3.4. Results. - 4. Corpus-based analysis. - 4.1. Results. - 5. Contextual preferences for *disease/illness.* - 5.1. Results. - 6. Conclusions. - 7. References.

1. Abstract

It often happens that terms defined by dictionaries as synonyms have different patterns of use. An example of this phenomenon is represented by the terminological pair *disease/illness* in medical discourse. The behaviour of these commonly used terms will be an object of discussion in this study.

The work is divided into two main sections. The first part takes into account the information included in conventional dictionaries, both bilingual and monolingual, generic and specialized, in order to see if the terms in question are described as perfectly interchangeable or as having different uses depending on the context. Some dictionaries highlight the use, both specific and generic, respectively of *disease* and *illness*, but limited information about frequency, collocational and contextual preferences is provided. The second part of the work is represented by two corpus-based studies. In the first corpus, made up of 30 research article texts, the frequency of *disease* and *illness*, as well as collocates and patterns of use, are investigated. In the second additional corpus, made up of other 20 research articles texts, the aim of the analysis is to find out if an association exists between the use of *disease* and contexts where *specific* pathological conditions are dealt with, and between *illness* and contexts having pathological conditions *in general* as a subject.

The work intends to draw attention to the enormous resources that corpus linguistics can offer to lexicographic research.

^{*} Researcher in English language and translation at the Faculty of Human Movement and Sports Sciences of the University of Palermo.

2. Introduction

Corpus-based linguistic investigations have achieved growing importance over the last few decades. Corpora allow researchers to observe and describe linguistic phenomena in a way that was not possible in the past with traditional approaches to language. Thanks to the extraordinary advances in information science technology and to the use of sophisticated electronic tools for linguistic analysis, today large amounts of authentic data are investigated quickly and accurately by means of a quantitative approach. In this way linguists are offered the possibility to make generalizations about several aspects of language which are not based upon personal intuitions. Moreover, the capability of modern softwares to retrieve and study words, phrases or even large pieces of text in context and not in isolation has enriched linguistic research with information about language use and grammatical behaviour of words with unprecedented results.

Corpora represent a very important source of information for learners, who, however, most times turn to the consultation of printed dictionaries. These are of course indispensable tools for language learning, but also have several important shortcomings. Bowker and Pearson (2002:15) highlight some of the main limitations of printed dictionaries in helping users with specialized language. In addition to the fact that dictionaries "go out of date very quickly", as by the time a dictionary is compiled and printed a language has acquired new terms, one of the most severe limitations is the size of dictionaries. Since many users prefer to have easy-to carry dictionaries, most books contain limited information about terms, and the information lexicographers choose to include does not always meet LSP users' needs. Moreover, Bowker and Pearson (2002:16) draw attention to one of the most critical features of printed dictionaries, that is the fact that they are not fulfilling in giving information about how terms are used in context. Printed dictionaries, in fact, generally provide the meanings of terms but do not say anything about collocates, that is other words which often associate with those which are the object of investigation.

Given the importance of contextualizing words and providing information about their usage, lexicographic research has recently paid more attention to corpora analysis for dictionary project purposes.¹

Biber et al. (1998:24) identify six major types of research questions in corpus-based lexicographic investigations:

- 1. meanings associated with a particular word (traditional focus of lexicography);
- 2. the frequency of a word relative to other related words;
- 3. non-linguistic association patterns of a particular word (registers, historical periods, dialects, and so forth);

¹ Descriptions of how corpora are used in lexicography today are offered by Pulcini (2008), and Hunston (2002: 96-108).

- 4. common collocates of a particular word, and distribution of collocational sequences across registers;
- 5. distribution of the senses and uses of a word;
- 6. use of seemingly synonymous words and their distribution in different ways.

As far as the last issue is concerned, Biber et al. (1998:24) state:

Languages have many words that are similar, and dictionary definitions often characterize such words as identical or "synonymous" in meaning. The patterns of use for these words, however, are often very different. Investigating the use and distribution of synonyms in a corpus allows us to determine their contextual preferences, associated with other collocates or associated with register differences.

In this study the pattern of use of the two medical terms *disease* and *illness* will be an object of investigation.

3. The case of disease and illness

Disease and illness are certainly among the most frequently used terms in medical language. Many dictionaries consider them as synonyms, even though *disease* is often described as having a specific use and *illness* a more general one. In this study several dictionaries were consulted in order to find out whether an actual difference between the two terms could emerge from definitions and from possible usage examples. To this purpose bilingual generic and specialized dictionaries, and monolingual generic and specialized dictionaries were consulted. Generic bilingual dictionaries were consulted in order to see what Italian equivalents were registered and whether any examples contextualizing both terms could be found. More in general, both generic bilingual and monolingual dictionaries were taken into account in this study basically because disease and illness are terms which do not belong only to medical discourse, but to common language as well. As far as specialized dictionaries are concerned, these were consulted to find out if any difference of use could be straightforwardly deduced from the definitions of the terms in question, as well as, of course, from any related examples.

3.1. Consulting generic bilingual dictionaries

The following is what can be read with regard to *disease* and *illness* in the Garzanti (2002) and Picchi (1999) bilingual dictionaries:

| Disease malattia |
|---|
| Occupational disease, malattia |
| professionale. |
| sin. Illness |
| Confrontiamo disease e illness. |
| Entrambi i termini hanno lo stesso |
| significato e spesso sono usati uno per |
| l'altro. Si può dire però che disease indichi |
| la malattia specifica che si prende, si |
| trasmette, si cura ed è oggetto di studio da |
| parte dei medici: This plant suffers from a |
| rare disease, Questa pianta soffre di una |
| rara malattia; infectious disease, malattia |
| infettiva; skin disease, malattia della pelle. |
| Illness è più generico e in genere è riferito a |
| persone: She died after a long painful |
| illness, Morì dopo una lunga e penosa |
| malattia. ² |
| Illness malattia, malanno. |
| sin. disease |

Table 1. Extract from Garzanti (2002)

| Disease | | |
|------------|---|-------------|
| 1. malatti | a, morbo | |
| Ex: Alzh | eimer's disea | se, morbo o |
| malattia d | li Alzheimer. | |
| 2. malatti | a, morbo, mal | e, vizio |
| Illness | malattia, | infermità, |
| malanno, | disturbo. | |
| | $\mathbf{D} \leftarrow \mathbf{C} = \mathbf{D}$ | 11 (1000) |

Table 2. Extract from Picchi (1999)

As can be seen from table 1, the only Italian equivalent which is registered for *disease* is *malattia*, unlike what happens in table 2, where *morbo* can also be found with regard to eponyms (*Alzheimer's disease*). Moreover, in table 1 it is first said that *illness* and *disease* have the same meaning, and then that the terms are used in a different way, *disease* having a more specific meaning (it is used to refer to conditions which are caught, transmitted, treated, and which are also an object of attention by physicians), while *illness* is more generic and is usually used when dealing with people (*she died after a long painful illness*).

Finally, looking once again at table 2, *male* and *vizio* are registered as additional Italian equivalents for *disease*, but there is no trace of them in table 1.

² The following is a comparison between *illness* and *disease*. The two terms have the same meaning and are often used interchangeably. Yet, *disease* refers to a specific condition which is caught, transmitted, treated, and which is studied by physicians. Ex: *This plant suffers from a rare disease; infectious disease; skin disease. Illness* is more generic and is usually used with reference to people: *She died after a long painful illness*; Morì dopo una lunga e penosa malattia. (My own translation).

As far as *illness* is concerned, the Italian equivalents registered in table 1 are *malattia* and *malanno*, which can also be found in table 2 next to *infermità* and *disturbo*.

The definitions reported above do not, however, cast sufficient light on the difference of use between *illness* and *disease*, firstly because the only information which can be inferred is that *disease* refers to a specific pathological condition, and *illness* to a generic pathological one which may not necessarily coincide with a sheer condition of 'poor health', and secondly because there are not enough examples clarifying this potential difference.

3.2. Consulting specialized monolingual dictionaries

Dorland's Illustrated Medical Dictionary (2007) defines *disease* "any deviation from or interruption of the normal structure or function of a part, organ, or system of the body as manifested by characteristic symptoms and signs; the aetiology, pathology, and prognosis may be known or unknown". At the entry *illness*, the only information provided is that the term is a synonym of *disease*. Two examples for *illness* are registered, *folk illness* and *psychosomatic illness* (or *somatoform disorder*). The first is "a condition in which a person has symptoms that are not identifiable with usual modern categories of disease", and whose name is explained with the fact that in many societies this condition and other which are similar "are felt to be treatable by folk healers using techniques such as prayer, rituals, and laying on of hands"; the second is a type of condition belonging to the group of mental disorders, "in which physical symptoms suggest the presence of a medical disorder but are not fully explained by a general medical condition".³

In Medline Plus Medical Dictionary:

(www.nlm.nih.gov/medlineplus/mplusdictionary.html), one of the most consulted dictionaries on the Internet, *disease* and *illness* are defined as follows:

disease an impairment of the normal state of the living animal or plant body or one of its parts that interrupts or modifies the performance of the vital functions, is typically manifested by distinguishing signs and symptoms, and is a response to environmental factors (as malnutrition, industrial hazards, or climate), to specific infective agents (as worms, bacteria, or viruses), to inherent defects of the organism (as genetic anomalies), or to combinations of these factors: syn. illness, sickness; called also *morbus*

illness an unhealthy condition of body or mind: syn. sickness

The results of the investigation on *disease* and *illness* as carried out in the two specialized monolingual medical dictionaries mentioned so far do not prove to be fulfilling enough to the purpose of this work. Neither usage examples nor any explanation of the possible semantic difference between the two terms are offered. Undoubtedly, the terms chosen for the analysis are so frequently used in

³ For investigations about the use of *disorder* in the sense of *illness*, see Gavioli (2000).

medical language that it would not be feasible for dictionaries to provide a sufficiently significant number of examples to allow deducing the possible semantic preferences of *disease* and *illness* in medical language. However, the need for a deeper investigation on the issue in question is felt as necessary, especially when running into such definitions as that provided by the online encyclopaedia *wikipedia* (www.wikipedia.org), where, with regard to *illness* (also compared to *sickness*), we read:

Illness and sickness are generally used as synonyms for disease. However, this term is occasionally used to refer specifically to the patient's personal experience of his or her disease. In this model, it is possible for a person to be diseased without being ill (to have an objectively definable, but asymptomatic, medical condition), and to be ill without being diseased (such as when a person perceives a normal experience as a medical condition, or medicalizes a non-disease situation in his or her life).

In "it is possible for a person to be diseased without being ill", and "to be ill without being diseased", the adjectives "diseased" and "ill" have a different semantic value, that is diseased refers to a person who is actually in a state of 'lack of health', that is he or she is in a medical condition which is "objectively definable", while the term *ill* can also be used to designate a person who considers himself or herself in a condition of 'lack of health', but is not actually so. Thus, *ill* has a subjective value, that is it can be used to refer to *how* a person perceives his or her state of being, while *diseased* has an objective value; it is used to refer to *actual* states of pathological conditions, free from possible personal perceptions.

3.3. Consulting other dictionaries

To the purpose of this study, and in order to have a larger view of definitions and examples offered by dictionaries, the generic monolingual *Oxford English Dictionary* (2000), *Merriam-Webster Online* (www.m-w.com), and the bilingual specialized *Gould-Chiampo* (1998) dictionary were also consulted. The following is what was found in these works:

a. Oxford

A **disease** is a particular illness with a name, or an illness which affects a particular part of the body:

- Measles is the most devastating of all the major childhood diseases.
- A healthy diet and regular exercise can help prevent heart disease.

Illness is a general word for a period of not being in good health:

- He died unexpectedly after a short illness
- The doctor asked whether she had a history of any serious illness

b. Merriam-Webster Online

Disease an impairment of the normal state of the living animal or plant body or one of its parts that interrupts or modifies the performance of the vital functions, is typically manifested by distinguishing signs and symptoms, and is a response to environmental factors (as malnutrition, industrial hazards, or climate), to specific

infective agents (as worms, bacteria, or viruses), to inherent defects of the organism (as genetic anomalies), or to combinations of these factors: syn. illness, sickness - called also *morbus*

Illness an unhealthy condition of body or mind: syn. sickness

c. Gould-Chiampo

Disease *n*. **malattia.** The failure of the adaptive mechanisms of an organism to counteract adequately the stimuli or stresses to which it is subject, resulting in a disturbance in function or structure of any part, organ, or system of the body. A response to injury; sickness or illness.

Illness *n*. malattia; indisposizione. **1**. The state of being ill or sick. **2**. A malady, sickness, disease, disorder.

3.4. Results

On the basis of the data obtained from the consultation of the dictionaries mentioned so far about the use of *disease* and *illness* it seems that the former is likely to occur in contexts dealing with pathological conditions which affect a specific part of the body (examples mentioned above are "heart disease", "eye disease", "kidney disease" or "disease of the kidney", while the latter has a more generic use (e.g. "various illnesses", "short illness", "because of illness"), that is it is used in contexts where there is no reference to a *specific* pathological condition.

4. Corpus-based analysis

At this stage of the research, the need for an evidence-based analysis of authentic texts was felt, that is a corpus-based linguistic investigation in order to see how the terms chosen for this study are actually used in authentic medical English language and what "contextual preferences" they show to have. To this purpose, a corpus of 109,822 words from 30 English texts was collected. All texts are research articles drawn from the BMJ (British Medical Journal) online, covering the period from 2007 to 2009.⁴ The average number of words per text is 2,929, ranging from 638 words to 5,220. All sections were inspected in the texts, except authors' names, titles, abstracts, acknowledgements, and references. The corpus covers a wide number of different medical branches, including cardiology, dermatology, neurology, internal medicine, intensive care, oncology, orthopaedics and rheumatology, surgery, obstetrics, and infectious diseases. The linguistic tool used for the analysis was WordSmith (version 5.0), a software producing concordances in the Key Word in Context (KWIC) format.

The study aimed at the investigation of the two main following issues:

- a) frequency of *disease* and *illness* in the semantic areas explored;
- b) collocates associated with *disease* and *illness*.

⁴ The English texts selected for this study were drawn from the following issues: 334 (May 2007), 335 (June, July, and August 2007), 336 (February, March, and April 2008), 338 (January, February, and May 2009), and 339 (October 2009).

Disease showed 194 occurrences (accounting for 85.5%), while *illness* showed 33 occurrences (accounting for 14.5%).⁵

Disease showed a tendency to collocate mostly with nouns and adjectives respectively designating and pertaining to organs. "Heart" is the top collocate noun, with 24 occurrences, followed by "cuff" with 15 occurrences (although concentrated in a single text), "kidney" with 9 occurrences, "bowel" with 5 occurrences, and "liver" with 3 occurrences.⁶

The top collocate adjective is "meningococcal" with 21 occurrences (although concentrated in a single text), followed by "cardiovascular" with 7 occurrences, "vascular" and "chronic" with equal number (6) of occurrences, "pulmonary" and "histological" with equal (5) number of occurrences, and "cerebrovascular" with 4 occurrences.

| Nouns | Occurrences |
|--------|-------------|
| Heart | 24 |
| Cuff | 15 |
| Kidney | 9 |
| Bowel | 5 |
| Grade | 5 |
| Liver | 3 |
| Total | 61 |

Table 3. Left-collocate nouns of *disease* in the selected corpus from the BMJ (109,822 words).

| Adjectives | Occurrences |
|-----------------|-------------|
| Meningococcal | 21 |
| Cardiovascular | 7 |
| Chronic | 6 |
| Vascular | 0 |
| Pulmonary | 5 |
| Histological | 5 |
| Cerebrovascular | 4 |
| Total | 54 |

Table 4. Left-collocate adjectives of *disease* in the selected corpus from the BMJ (109,822 words).

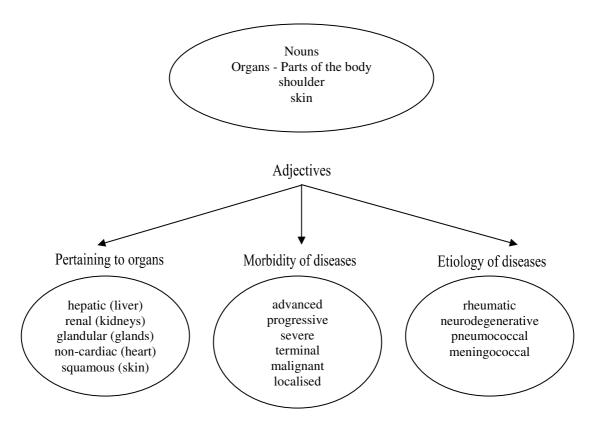
4.1. Results

The results showed that 31.4% of left-collocates of *disease* are nouns mainly representing names of organs. Nouns not representing names of organs are "grade", with 5 occurrences, which is synonym of "level" of disease, "reflux" (1 occurrence), which was not included in table 3, and "cuff", which although not being an organ, is a part of the body; 27.9% of left-collocates was represented by adjectives pertaining organs and tissues (e.g. "pulmonary" refers to lungs), except "chronic", which describes the course of a disease (or its rate of onset and

⁵ The concordances of *disease* and *illness* performed by WordSmith are fully shown in Appendix 1 of this work.

⁶ Nouns and adjectives with less than 3 occurrences were not included in table 3 and 4. They will be mentioned later in the text.

development); 21.7% is represented by articles and prepositions ("the", "a", "on", "for", "in", "to", etc.); 12.3% is made up of adjectives and (a few) nouns, which, although recurring most times only once, were considered important as they may contribute to giving information about the semantic preferences of *disease* in medical discourse. These terms were divided as follows:



The remaining terms are represented by the adjectives "future", "resultant" and "known", which owing to their non-specialized nature were defined here as 'conversational' terms, and by the adjectives "pilonidal", an anatomo-pathological definition, and "communicable", referring to the type of disease. The noun "reflux" and the names "Osgood-Schlatter" are, respectively, a physiopathological definition and an eponym.⁷

As far as the concordance of *illness* is concerned, the results showed that, firstly, this term mainly associates with adjectives (accounting for 60.7%)⁸, and, secondly, that these adjectives refer to different semantic areas. "Psychotic", for instance, describes the 'type' of illness, in this case with reference to the psychiatric area; "prodromal" and "unstable" are terms referring to the 'development' of a pathological condition; "medical" designates the classification

⁷ The remaining 6.7% is mainly represented by punctuation marks (full stops and commas), thus not giving any meaningful information about collocates of *disease*.

⁸ The remaining 39.3% is mainly represented by prepositions ("after", "of", "to", etc).

of diseases according to treatment, while "critical" and "severe" belong to the sphere of 'morbidity' of pathological conditions. Though the occurrences of *illness* are not, perhaps, numerically enough to allow drawing generalizations, nevertheless a more general use of the term seems to be suggested by its concordance in comparison with that of *disease*.

5. Contextual preferences for disease/illness

To complete the work done so far in this study, additional research was carried out in order to investigate if what was registered in dictionaries about the association of specific pathological conditions with *disease* rather than with *illness* could be confirmed by a corpus-based analysis. To this purpose, a different corpus was analysed.⁹ It is made up of 20 research article texts from the BMJ online, accounting for 95,288 words, and covering the period of time 2006-2010. The average number of words per text is 4,413, ranging from 2,365 to 6,461.¹⁰ The texts deal with several pathological conditions, including measles, pneumonia, diabetes, obesity, multiple sclerosis, cirrhosis, and many others.

In order to carry out this analysis, it was necessary to investigate larger strings of text than the short fragments of word sequences to the left and right of the chosen keyword in each line. More precisely, chunks of texts were inspected every time any pathological condition was mentioned.

5.1. Results

The investigation of the corpus described above showed that in texts dealing with specific conditions (that is those with a name) or with conditions affecting a specific part of the body the term used is *disease*, while *illness* most often occurs in contexts where the condition "of not being well" is dealt with, without reference to any specific pathological condition.¹¹ However, the analysis of the corpus revealed another important datum which was not found in any traditional dictionary consulted for this study, that is the use of *illness* with the same semantic value as *disease*; in other words, the use of *illness* as a synonym of *disease* also in contexts where there is reference to specific pathological conditions. More precisely, *illness*, which occurs 41 times in the corpus, shows 36 occurrences (accounting for 87.9%) revealing a general use of the term, and 5 occurrences (12.1%) where the term refers to specific pathological conditions ("myalgic encephalitis", "asthma", "diabetes", "heart failure" and "H1N1 influenza").¹²

⁹ In order to carry out this research it was thought it right to build another corpus with different design criteria which took into account texts about specific pathological conditions.
¹⁰ The English texts collected for this study were drawn from the following issues: 332 (January and March

¹⁰ The English texts collected for this study were drawn from the following issues: 332 (January and March 2006), 335 (August 2007), 336 (February and June 2008), 337 (July 2008), 339 (July, September, and December 2009), 340 (February and March 2010).

¹¹ Some text fragments from the concordance of *illness* in the second corpus are included in Appendix 2 (b).

 $^{^{12}}$ The texts fragments where these pathological conditions are mentioned in association with *illness* can be found in Appendix 2 (b), extracts 6 and 7.

A look at the entry *illness* in the corpus-based *Collins COBUILD English Language Dictionary* (2001) seems to confirm two uses of the term, one of which describes *illness* as a synonym of *disease*:

Illness 1 is the experience of being ill for a period of time. EG During his last illness we only saw him twice...People can recover from the symptoms of mental illness.

Illness 2 An illness is a particular disease that people can suffer from, such as cold, measles, or pneumonia. EG She died of a mysterious illness...Finally a doctor diagnosed the illness...cures for various illnesses.¹³

Collin's Dictionary of Medicine (2001), another corpus-based dictionary, also registers the two different uses of *illness*:

Illness (a) state of being ill, of not being well;

- His illness makes him very tired;
- Most of the children stayed away from school because of illness
 (b) type of disease;
- He is in hospital with an infectious tropical illness;
- Scarlet fever is no longer considered to be a very serious illness¹⁴

Disease, occurring 283 times, is always found in contexts where specific pathological conditions are dealt with ("diabetes", "psoriasis", "whooping cough", etc.).¹⁵

6. Conclusions

The analysis of the small medical corpora built for this work does not claim to give any conclusive data. The aim of the investigation was only that of identifying the collocational and contextual preferences of *disease* and *illness*, two very common medical terms which are often described as synonyms by dictionaries, though having different patterns of use. Dictionaries, in fact, though representing indispensable source of information about the meaning of words, do not generally provide enough usage examples and information about collocates or semantic preferences of terms. Not even information about the frequency of a given term is offered.

From the analysis of the first corpus it can be inferred that *disease* shows a clear tendency to (a) occur more frequently than *illness*, and (b) collocate with nouns and adjectives referring to organs and parts of the body. *Illness* shows a tendency to collocate almost exclusively with adjectives, with these belonging to

¹³ As far as *disease* is concerned, *Collins COBUILD English Language Dictionary* (2001) registers: **Disease** An illness in people, animals or plants which is caused by bacteria or infection, rather by an accident. **EG**...women and children ravaged by disease...I have a rare eye disease....infectious diseases.

¹⁴ Collin's Dictionary of Medicine (2001) defines disease "illness (of people, animals, plants, etc.) where the body functions abnormally";

She is suffering from a very serious disease of the kidneys or from a serious kidney disease;

He is a specialist in occupational diseases or in diseases which affect workers.

¹⁵ See Appendix 2 (a).

different semantic areas. None of the left collocates of *illness* refers to organs or parts of the body.

Another corpus-based analysis was carried out in a different corpus in order to investigate if what was registered in dictionaries about the use of *disease* in contexts dealing with specific pathological conditions could be confirmed. The analysis of the corpus revealed a clear prevalence of *disease* in such contexts, while *illness* was prevalent in more general contexts. However, the investigation of the corpus revealed the use of *illness*, though in a low percentage, as a synonym of *disease* also in contexts where specific pathological conditions were dealt with.

It can be stated that studies like the one carried out here can prove useful, for instance, from a translating point of view. In the case of *disease* and *illness*, if on the one hand both terms can be rendered into Italian as "malattia" (or often "morbo" in the case of *disease* used in eponyms), on the other one when translation is *into* English the semantic preferences and collocational patterns of either terms should be known.

Finally, with regard to the role and importance of dictionaries within ESP, it is worth saying that even though they are no doubt a valuable information source for learners to get definitions and meanings of terms, they should also provide information about frequency of terms, as well as about context, and give more room to usage examples. This entails two major requisites, the necessity for dictionaries to be corpus-based, and the need for them to have an online format. The latter would also allow entries to be continually enriched and updated with the latest research outcomes.

Examples of major corpus-based dictionaries (for common English) are *Collins COBUILD English Language Dictionary*, created after a project set up by John Sinclair, and *Longman Dictionary of Contemporary English*.

A bilingual English-Italian/Italian-English corpus-based specialized medical dictionary is still lacking. It would certainly enrich Italian lexicographic research within ESP studies and be a valuable aid for translating purposes as well.

7. References

1. Biber D., Conrad S., Reppen R., 1998, *Corpus Linguistics: Investigating Language Structure and Use*, Cambridge, Cambridge University Press.

2. Bowker L., Pearson J., 2002, *Working with Specialized Language*, London and New York, Routledge.

3. Collin P., 2001, Dictionary of Medicine, Peter Collin Publishing.

4. Garzanti - Petrini, 2002, *Dizionario di Inglese Inglese-Italiano/Italiano-Inglese*, Milano, Garzanti Linguistica.

5. Gavioli L., 2000, "Some Thoughts on the Problem of Representing ESP through Small Corpora", in Kettemann B. & Marko G. (eds.), *Teaching and Learning by Doing Corpus Analysis. Proceedings of the Fourth International Conference on Teaching and Language Corpora*, Graz 19-24 July, Amsterdam & New York, Rodopi, pp. 293-303 (11).

6. Hunston S., 2002, *Corpora in Applied Linguistics*, Cambridge, London, Cambridge Applied Linguistics.

7. Il Gould Chiampo, Dizionario Enciclopedico di Medicina Inglese-Italiano/Italiano-Inglese, Chiampo L. (a cura di), 1998, Bologna, Zanichelli/McGraw-Hill.

8. Martelli A, Virginia P. (eds), 2008, *Investigating English with Corpora*, Monza, Polimetrica International Scientific Publisher.

9. *Medline Plus Medical Dictionary:* (www.nlm.nih.gov/medlineplus/mplusdictionary.html),

10. Merriam-Webster Online Dictionary (www.m-w.com)

11. Newman Dorland, W.A., 2007, *Dorland's Illustrated Medical Dictionary*, Philadelphia, W.B. Saunders Company.

12. Picchi F., 2002, Grande Dizionario di Inglese, Milano, Hoepli.

13. Pulcini V., 2008, "Corpora and lexicography: the case of a dictionary of Anglicisms", in Martelli A. & Pulcini V. (eds.), *Investigating English with Corpora*, Monza, Polimetrica International Scientific Publisher, pp. 189-203.

14. Sinclair J., 2001, *Collins COBUILD English Language Dictionary*, Toronto, HarperCollins Canada.

15. Wikipedia (www.wikipedia.org)

Appendix 1

Figure 1. Concordance of *disease* in the corpus from the BMJ (109,822 words), sorted by the first word to the left.

N Concordance sex]+[0.8618xcerebrovascular^ disease]+[3.2636xcardiogenic 2 the Kujala Patellofemoral[^] Scale, a disease specific validated disability scale з Introduction Pilonidal sinus is a disease that most commonly arises in 4 If a participant reported having had a disease diagnosed^ by a doctor for six 5 Dukes? A and B and "advanced disease" constitutes tumours infiltrating 6 threaten immunisation[^] targets and disease control should this concern 7 The association of preterm birth and disease has not been described? 8 hypertension, coronary[^] artery disease, percutaneous transluminal 9 diabetes, and coronary artery disease^ were identified as risk factors 10 less likely to have coronary artery disease (P = 0.04) or liver^ disease (P = 11 intraepithelial[^] neoplasia grade 2, as disease of this severity or greater was[^] 12 concerns about autism and bowel disease despite a balanced 13 link between MMR vaccination, bowel disease, ^ and autism (50% considered it concerns about autism and bowel disease.^ The most important perceived 14 15 concerns about^ autism and bowel disease (78%, /v/ 27%), and fear of instruments: the inflammatory bowel/ disease guestionnaire (UKIBDQ), 16 17 previous^ two weeks, any non-cardiac disease likely to limit survival^ to two 18 to1.51) and^ death from cardiovascular disease or hospital admission for^ heart 19 based studies of cardiovascular[^] disease in British towns representing all 20 with pre-existing cardiovascular disease, ^42 more research is ^ needed to 21 and determinants^ in cardiovascular disease (MONICA) criteria,^21 using 22 of their risk of cardiovascular[^] disease.[^]9 Only one qualitative study to be aware of the risk of cardiovascular disease associated^ with impaired 23 24 developing diabetes and cardiovascular disease.^ The lack of anxiety associated 25 for dialysis status, cerebrovascular disease, peripheral vascular disease. 26 ?80 years, female sex, cerebrovascular disease,^ cardiogenic shock, urgent 27 for dialysis status, cerebrovascular disease, peripheral vascular disease, 28 dialysis^ status, cerebrovascular disease, peripheral vascular disease,^ 29 be related to diagnosis[^] of a chronic disease (or the treatment in terms of in analysis*?Age, sex, modified chronic disease^ score, prescription drugs 30 account demographic^ factors, chronic disease, psychosocial risk factors in the 31 32 models for age,^ education, chronic disease, family status, health behaviour,^ 33 him or her^ as having a chronic disease.^ / Health behaviour/ We divided 34 sex, education, family status, chronic disease, and health^ behaviour.^15 The 35 illness.^ Consultants in communicable disease control, clinicians, the^ national renal, and retinal complications. Disease onset^ may occur up to 12 36 37 Non-operative treatment for rotator cuff disease primarily consists^ of active million, to refer patients with rotator cuff disease^ to the outpatient clinic of the and mechanisms of pain in rotator cuff disease^ are not known.^3 38 39 40 of steroid^ injections for rotator cuff disease is unconvincing. Conclusions^ of 41 impingement syndrome^ or rotator cuff disease is the most frequent 42 in patients^ with chronic rotator cuff disease. They found a moderate corticosteroid^ injections for rotator cuff disease. 9 ^11 Alvarez et al compared^ 43 corticosteroid injections^ for rotator cuff disease. Limited evidence exists for 44 45 strategy for patients with rotator cuff disease. Better outcome^ in terms of 46 corticosteroid injection in rotator cuff disease. ^ *Comparison with existing by the duration of rotator cuff disease. The favourable result^ of 47 48 short term improvement in rotator cuff disease. Ultrasound^ guided injection of bursa in patients with rotator cuff disease.^ We used a double blind 49 50 injections in patients with rotator cuff disease, we did a randomised^ controlled

SEZIONE 1

| 51 | in partients with rotator cuff disc | ÷ |
|-----|---|---|
| 52 | | ease specific° worry was measured |
| 53 | It addressed the question: does dis | |
| 54 | were used: "pilonidal" sinus" ,"fistula" ,"dis | |
| 55 | improvements in health, risk factors for dis | ease, or even overall levels of |
| 56 | fitness, other risk factors ^a for dis | ease, health, and wellbeing; evidence |
| 57 | neither necessary nor sufficient" for dis | ease. Its significance was unchanged |
| 58 | | ease susceptibility Intention to |
| 59 | or undifferentiated by age, gender, °dis | ease or organ. This includes |
| 60 | detection of histological ^a glandular dis | ease, although this result is limited by |
| 61 | more cases of histological ^a high grade dis | ease than did conventional cytology |
| 62 | more cases" of histological high grade dis | ease than did conventional cytology^ |
| 63 | more cases of histological* high grade dis | ease per 1000 women screened than |
| 64 | cases (21%) of histological* high grade dise | ease were detected (table 4Go). The |
| 65 | more cases of histological* high grade dis | ease and required 170 (380 minus |
| 66 | in adults with ischaemic heart ^o disc | ease. ^w10 None found a significant |
| 67 | service framework for coronary heart ^o disc | ease. ^1 Recently it has been |
| 68 | (adults with ischaemic heart dis | ease° w10 or type 2 diabetes° w23). |
| 69 | congestive heart failure, ^ valvular heart dis | ease, arteriosclerosis and |
| 70 | reductions in the risk of coronary heart dis | ease of up to 75% or more. ^2 Most |
| 71 | Secondary prevention of coronary heart dis | ease is an important component of |
| 72 | after admission, who had valvular heart dis | ease (defined as having inpatient |
| 73 | on diagnoses of coronary heart dis | ease. We used questionnaires in |
| 74 | intervention for patients with heart dis | ease was effective, ^w10 but studies |
| 75 | • | ease cases and attenuated by death |
| 76 | - +F | ease, heart failure, cholesterol |
| 77 | use ^a among prevalent coronary heart dis | |
| 78 | • • | ease. Self reported use of medication |
| 79 | • | ease, type 2 diabetes, and cancer of |
| 80 | | ease in older patients, particularly by |
| 81 | ,, | ease, pulmonary embolism, ^ |
| 82 | | ease, paintonary embolism, ease, heart failure, mean arterial |
| 83 | ,,, | ease was defined [°] by using the |
| 84 | | ease. ^20 Coronary heart disease was |
| 85 | | ease, heart failure, cholesterol |
| 86 | | ease, and use of psychotropic drugs |
| 87 | ,,, | ease, and use of psychotropic unugs ease, heart failure, cholesterol |
| 88 | | ease, and use of psychotropic drugs. ^ |
| 89 | | |
| | | , , |
| 90 | | ease, ^ anaemia *Agents |
| 91 | · · · · · · · · · · · · · · · · · · · | ease of cervical ^a intraepithelial |
| 92 | ••••• | ease than does conventional |
| 93 | •••• | ease by 1.3 cases per 1000 women |
| 94 | * * • • | ease detected* by imager read |
| 95 | * * • | ease detected by imager read |
| 96 | York Heart Association class For II dis | · / · |
| 97 | towards older teenagers ⁴ 4 and a rise in dis- | * 1 |
| 98 | York Heart Association class III or IV disc | |
| 99 | trials in patients with chronic kidney dis | , , |
| 100 | patients with advanced [*] chronic kidney dis | ease, we consider that renal |
| | | |

| immune mediated, polycysto kidney disease, other) "age at transplantation, in patients" with dronic kidney disease controlled" trials in chronic kidney disease controlled" trials in chronic kidney disease patients with advanced chronic kidney disease tage 3 or 4 were" randomly because a risk from" a known disease marey disease (P = 0.0 di) or liver" disease, "acmentia or cognitive antery disease (P = 0.0 di) or liver" disease, "acmentia or cognitive marey disease (P = 0.0 di) or liver" disease, "acmentia or cognitive antery disease (P = 0.0 di) or liver" disease, "acmentia or cognitive treatment or radiotherapy for maingrant disease, for each disease, "acme chronic andiac or finct of risk factor for meningococcal disease, and vere-estimate" protective in the model, with meningococcal disease in tenjalerst" are poorty the indence" of meningococcal disease in tenjalers" are poorty the indence" of meningococcal disease sa life threatening condition, " twith higher risk of meningococcal disease thigh season of disease in ubid season of meningococcal disease thigh season of the peak in meningococcal disease thigh season of disease thigh season of disease thigh season of disease thigh season of meningococcal disease thigh season of meningococcal disease thigh season of meningococcal disease thigh season of menin | | | | |
|---|-----|---|----------|---|
| 103 controlled" trials in chronic kidney disease and dialysis patients and 104 003 patients" with dwaned chronic kidney disease stage 4 for three years. "3 105 patients with advaneed chronic kidney disease for a higher mortality or more 105 patients with dwaneed chronic kidney disease stage 3 or 4 were" randomly 109 because a risk form" a known because are stage are form an anothing or coding. there registry was established by 101 attern disease, provides an evidential therease, here disease, "dementia or cognitive" 101 attern disease, provides an evidential disease, provides an evidential "severe chronic cardiac or 111 disease, provide and meningeococal disease, our work" provides an evidential disease 112 revision) for coding, in which "nonligeococal disease our work" provides an evidential 113 treatment or radiotherapy for meningeococal disease our work" provides an evidential 113 interded, with meningeococal disease our work" provides an evidential 114 interded, with meningeococal disease interagers" are poorly 115 refeteding frame meningeococal </td <td>101</td> <td>immune mediated, polycystic kidney</td> <td>disease,</td> <td>other) ,° age at transplantation,</td> | 101 | immune mediated, polycystic kidney | disease, | other) ,° age at transplantation, |
| 104 603 patients' with chronic kidney disease stage 4 for three years. "3 105 patients with advanced chronic kidney' disease. theraper advanced chronic kidney' disease. theraper advanced chronic kidney' disease. 107 1432 patients with chronic kidney' disease. theraper advanced chronic kidney' disease. theraper advanced chronic kidney' disease. theraper advanced chronic kidney' disease. 108 because a risk from's known disease. theraper advanced chronic disease. theraper advanced chronic disease. there registry was established by 109 haerontrage, renal disease. filesase. theraper advanced chronic disease. theraper advanced chronic cardiac or 111 treatment or radiotherapy br malignant disease. "severe chronic cardiac or 119 treatment or radiotherapy br malignococal disease. mostitutes Dules? A and B 119 the indedence" of meningcooccal disease in the angers" are poorly 119 the indedence" of meningcooccal disease in adolescence to examine 120 in the model, with meningcooccal disease in adolescence to examine 121 cohont study of meningcooccal disease in adolescence to examine 122 htroductin | | | disease | , |
| 105 patients with advanced chronic kidney' disease fund a higher mortality or more 106 patients with advanced chronic kidney' disease fund a higher mortality or more 107 1432 patients with chronic kidney' disease fund a higher mortality or more 108 because a risk from' a known disease fund a higher mortality or more 109 hearontrage, renal disease, liver disease "dementia or cognitive 101 artery disease (P = 0.04) or liver' disease "constitutes Dukes? A and B 111 disease, psychiatric disorders, "liver disease, "severe chronic cardiae or 112 treatment or radiotherapy to malignant disease are protecting' against meningcooccal disease or work' provides an evidential 119 the model, with meningcooccal disease in dolescence ' or examine isease 119 the model, with meningcooccal disease in dolescence ' or examine isease 120 in the model, with meningcooccal disease in dolescence ' or examine 121 cohont study of meningcooccal disease in dolescence ' or examine 122 htroduction Inmisococcal disease in dolescence' and the United 12 | | , | disease | and dialysis patients and |
| 108 patients with advanced chronic kidney' disease The registry was established by 107 1432 patients with chronic kidney' disease stage 3 or 4 were' randomity 109 because a risk form's known disease "dementia or cognitive "dementia or cognitive 100 artery disease, (P = 0. 04) or liver' disease, '' events or coding, in which 'localised disease "dementia or cognitive 111 disease, psychiatric disorders, 'liver disease, '' events or boronic cardiac or a sere thronic disease in the readers'' are poorty 118 effect of risk factors for meningcococal disease in adolescence to examine 121 eohont study of meningcococal disease in adolescence to examine 122 htroduction Invasive meningcococal disease in adolescence to examine 123 were at higher risk of meningcococal disease in adolescents' and a specific 124 with higher risk of meningcococal | | 603 patient <i>s</i> * with chronic <mark>kidney</mark> | disease | stage 4 for three years. ^3 |
| 107 1432 patients with chronic kidney disease stage 3 or 4 were " randomly 108 because a risk from" a Innoun disease, " dementia or cognitive 109 attery disease (P = 0.04) or live" disease, " dementia or cognitive 110 attery disease (P = 0.04) or live" disease, " dementia or cognitive 111 disease, psychiatrio disorders, " liver disease, " evere chronic cardiac or 112 revision) for coding, in which "localised disease, " as reported elsewhere. "23 The 119 inportant risk factor for meningcooccal disease on vork" provides an evidential 110 in protecting" against meningcooccal disease in tenagers" are poorly 119 The incidence" or meningcooccal disease in adolescence to examine 120 htroduction Imasive meningcooccal disease in adolescence to examine 121 orboot study of meningcooccal disease in sub-shana htroduction Imasive meningcooccal 121 otrisk factors for meningcooccal disease in sub-shana htria 122 htroduction Imasive meningcooccal disease in sub-shana htria 123 were at higher risk of meningcooccal disease in sub-shana | | · · · · · · · · · · · · · · · · · · · | disease* | • <i>i</i> |
| because a risk from a known disease haerrorthage, renal disease, liver disease, artery disease (P = 0. 04) or liver disease, revision) for coding, in which "localised treatment or radiotherapy for malignant linked to lower risk" of meningococcal effect of risk factors for meningococcal fine reasons for the peak in meningococcal disease, norther peaks of meningococcal disease fine reasons for the peak in meningococcal disease fine the rodel, with meningococcal disease fine there are higher fisk of meningococcal disease fine there are higher fisk of meningococcal disease fine there are higher fisk of meningococcal disease fisk factors for mening | | F, | disease. | The registry was established by |
| haemorthage, renal disease, liver disease, artery disease (P = 0. 04) or liver' disease, revision) for coding, in which 'localised disease' in terustric disease, medical "constitutes Dukes? A and B "severe chronic cardiac or as reported elsewhere. "23 The our work' provides an evidential and over-estimate' protective disease, in protecting' against meningococcal in protecting' against meningococcal in the model, with meningococcal in the model, with meningococcal disease in tenagers' are poorly in England' 2 and the United high season defined as 70 or as like threatening condition, " the tenagers' are poorly in tadelescence to examine is a life threatening condition, " than poople in employment. were history of preceding in sub-Saharan Affica or infuenza. The preceders' in adelescents' and a specific in tenagers' aged 15:19. Of 'Specimental and clinical "Outcome measures' Primary and economic productivity of acses mix, 'and list size have between' groups in our study. ' desput evidence of dise | 107 | · · · · · · | disease | stage 3 or 4 were [*] randomly |
| 110artery disease (P = 0. 04) or liver' disease, disease, psychiatro disorders, "liver' disease, revision) for coding, in which "localised disease, in portant risk factor for meningcooccal disease, in portant risk factors for meningcooccal disease, in portant risk factors for meningcooccal disease, in protecting" against meningcooccal disease, in protecting" against meningcooccal disease, in protecting" against meningcooccal disease, in protecting" against meningcooccal disease disease, in the model, with meningcooccal disease dis advectors for meningcooccal | 108 | because a risk from° a known | disease | may be more acceptable than a |
| 111disease, psychiatricdisorders, "liverdisease, revision) for coding, in which "localised disease, "constitutesDukes? A and B112revision) for coding, in which "localised treatment or radiotherapy for malignant linked to lower risk" of meningococcal of risk factors for meningococcal in portecting" against meningococcal in the model, with meningococcal of meningococcal disease lineases"severe chronic cardiac or as reported elsewhere. "23 The with high season defined as 70 or in adoesence to examine is a life threatening condition, " thigh season defined as 70 or in adoesence to examine is a life threatening condition, " thigh season of meningococcal disease thigh season defined as 70 or in adoesence to examine is a life threatening condition, " thigh season of meningococcal disease thigh season of meningococcal disease to risk factors for meningococcal | 109 | haemorrhage, renal disease, <mark>liver</mark> | disease, | ° dementia or cognitive |
| 112revision)for coding, in which "localised treatment or radictherapy for malignant linked to lower risk" of meningcoccal linked to lower risk" of meningcoccal disease, in protant risk factor for meningcoccal disease reasons for the peak in meningcoccal disease in the model, with meningcoccal disease high season defined as 70 or disease high season of meningcoccal disease high season of meningcoccal disease hi | 110 | artery disease (P = 0, 04) or liver* | disease | (P = 0. 02) (table 1). According |
| 113treatmentor radiotherapyfor malignantdisease, (sease, as reported elsewhere. "23 The114linked to lower fisk" of meningococcaldisease, (sease, in protecting" against meningococcaldisease, (sease, and over-estimate" protective116effect of fisk factors for meningococcaldisease, (sease, <br< td=""><td>111</td><td>disease, psychiatric disorders, °liver</td><td>disease,</td><td>rheumatic disease, medical</td></br<> | 111 | disease, psychiatric disorders, °liver | disease, | rheumatic disease, medical |
| linked to lower risk⁶ of meningooocal disease, as reported elsewhere. *23 The important risk factor for meningooocal disease, and over-estimate⁶ protective work⁶ provides an evidential and over-estimate⁶ protective work⁶ provides and over-estimate⁶ protective work⁶ provides and over-estimate⁶ protective work⁶ provides and provides and over-estimate⁶ protective work⁶ provides and over-estimate⁶ protective work⁶ provides and provides and over-estimate⁶ protective work⁶ provides and provides and provides and over-estimate⁶ protective work⁶ provides and provides an | 112 | revision) for coding, in which "localised | disease" | ° constitutes Dukes? A and B |
| 115importantrisk factorformeningococcaldisease, diseaseour work" provides an evidential and over-estimate" protective with increasing age as in tenagers" are poorly in England" 2 and the United high season defined as 70 or in adolescence to examine is a like threatening condult disease in adolescence to examine is a like threatening condult disease were at higher risk of meningococcal with higher" risk of meningococcal diseaseour work" provides an evidential and over-estimate" protective with increasing age as in tenagers" are poorly in England" 2 and the United high season defined as 70 or in adolescence to examine is a like threatening condult disease in adolescence to examine is a like threatening condult disease in sub-Saharan Africa or risk factors for meningococcal disease or risk factors for meningococcal disease or risk factors for meningococcal disease bis diseaseis likely that our cases were bisousion Risk factors for this likely that our cases were bisousion Risk factors for meningococcal disease bisousion Risk factors for meningococcal disease bisousion Risk factors for meningo | 113 | treatment or radiotherapy for malignant | disease, | ° severe chronic cardiac or |
| 116effect of risk factors for meningococcal in protecting* against meningococcal in protecting* against meningococcal in the measons for the peak in meningococcal disease in the model, with meningococcal disease htroduction Inwasive meningococcal disease were at higher* risk of meningococcal disease with higher* risk of meningococcal disease with higher* risk of meningococcal disease disease disease in adolescence to examine is a lift threatening of protecting in sub-Saharan Africa or inducta a history of preceding in sub-Saharan Africa or inducta. The precise* * Discussion Risk factors for meningococcal disease of risk factors for meningococcal disease of risk factors for meningococcal disease of risk factors for meningococcal disease of risk factors for meningococcal disease disease to risk* of exposure to meningococcal disease to risk* factors for meningococcal diseaseand over-estimate* in tengagers* age ordito, * than people in employment. were history of preceding induded a history of preceding induded a history of preceding inducted a history of preceding inducted a history of preceding inducted a history of preceding in sub-Saharan Africa in adolescents* and a specific in adolescents* and a specific in adolescents* differ* from those ligible* subjects were in adolescents differ* from those ligible* subjects were in tenagers* aged 15-19. Of * Discussion of mening concal disease ho differences existed in duration of disease specific epidemiological investigation of disease specific epidemiological investigation of disease specific epidemiological investigation of disease specific measures* HADS score, severity of disease or both). Laboratory confirmation of disease study will be the long* term rates of disease stu | 114 | linked to lower risk ^a of meningococcal | disease, | as reported elsewhere. "23 The |
| 117in protecting* reasons for the peak in meningococcal in the model, with meningococcal ochoit study of* meningococcal in the model, with meningococcal ochoit study of* meningococcal disease in teenagers* are poorly in England* 2 and the United high season defined as 70 or in adolescence to examine is a life threatening condition, * than people in employment. were history of preceding induded a history of preceding in adolescents* and a specific in adolescents* and a specific i | 115 | important risk factor for meningococcal | disease, | our work^ provides an evidential |
| 118reasons for the peak in meningococcal in the model, with meningococcal in the model, with meningococcal in the model, with meningococcal disease throduction Invasive meningococcal were at higher risk of meningococcal mix factors for meningococcal diseasein teenagers" are poorly in England" 2 and the United high season defined as 70 or in adolescence to examine is a life threatening condition, " usere history of preceding insub-Saharan Africa or finisk factors for meningococcal disease insub-Saharan Africa disease insub-Saharan Africa or finisk factors for meningococcal disease insub-Saharan Africa disease insub-Saharan Africa or finisk factors for meningococcal disease insub-Saharan Africa disease insub-Saharan Africa or sak factors for meningococcal disease insub-Saharan Africa disease insub-Saharan Africa disease in adolescents disease in adolescent teat in teenagers aged 15-19 | 116 | effect of risk factors for meningococcal | disease | and over-estimate [*] protective |
| 119The incidence* of meningococcal in the model, with meningococcal ochort study of meningococcal throduction Invasive meningococcal were at higher risk of meningococcal of meningococcaldisease disease disease disease disease diseasein England* 2 and the United high season defined as 70 or in adolescence to examine is a life threatening condition, * than people in employment. were at higher risk of meningococcal disease in sub-Saharan Africa or infuenza. The precise* or infuenza. The precise* or infuenza. The precise* or infuenza. The precise* tor high season of meningococcal disease or insk factors for meningococcal disease to risk* of exposure to meningococcal disease to risk* factors for meningococcal disease to risk* factors for meningococcal disease to risk* factors for meningococcal disease to nisk* factors for meningococcal disease to on risk factors for neurodegenerative disease to on risk factors for neurodegenerative of mannose-binding* lectin and risk of disease to ra genetic model in which severity of disease specific epiderriological investigation of disease to robt). Laboratory confirmation of disease to robt). Laboratory confirmation of disease to robt). Laboratory confirmation of disease to robt' used will be long term rates of disease to robt' used will be long term rates of disease. to robt' used will be long term rates of disease to robt' used will be long term rates of disease to robt' used will be long term rates of disease to robt' used will be long term rates of disease to robt' use | 117 | in protecting* against meningcooccal | disease | with increasing age as |
| 120in the model, with meningococcal cohot study of meningococcal httroduction httroduction linasive meningococcal were at higher risk of meningococcal diseasedisease disease disease disease disease high season defined as 70 or in adolescence to examine is a life threatening condition, ^ than people in employment. were history of preceding induded a history of preceding insub-Saharan Africa or infuenza. The precise * Discussion Risk factors for may predispose to meningococcal disease to risk' factors for meningococcal disease to risk' factors for meningococcal disease * Discussion disease * Discussion Risk factors for t is likely that our cases were . Further, findings concerning^* in adolescents* and a specific in adolescents* and a specific in adolescents* and a specific in adolescents* and a specific in adolescents* and a clinical * Dutorne measures* Primary and economic productivity of * Case mix, *and list size have between* groups in our study. * despite evidence that prevention, family experience of disease is independent* of genes risk' during the adolescent peak cannot* beinfing for a genetic model in which severity of disease rof a genetic model in which severity of disease rof the study will be long term rates of disease row the acolescent peak cannot* be inferred from the data was sought at the reference * Results at recruitment fom (APACHE II (acut physiology, The size (about 100 000 specific measures* '12 '13 * 14 | 118 | reasons for the peak in meningococcal | disease | in teenagers" are poorly |
| 121cohott study of meningcoccal htroductiondisease meningcoccalin adolescenceto examine is a life threatening condition, " than people in employment.123were at higher risk of meningcoccal uith higher' risk of meningcoccal isk factors for meningcoccal of risk factors for meningcoccal diseasedisease disease in duded a history of preceding in sub-Saharan or infuenza. The precise" "Discussion Risk factors for t is likely that our cases were . Further, findings concerning" in adolescents" and a specific in adolescents differ" from those124with kigher' nisk factors for meningcoccal to risk factors for meningcoccal to risk factors for meningcoccal diseasedisease disease or infuenza. The precise" "Discussion Risk factors for t is likely that our cases were . Further, findings concerning" in adolescents differ" from those133two winter peaks of meningcoccal presentation of recurrent or de now disease practice, including prevalence of disease specific endemiological investigation of disease specific model in which severity of disease specific endemiological investigation of disease specific endemiological investigation of disease specific endemiological investigation of disease specific endemiological investigation of disease study will be long term rates of disease study will be long term rates of disease a study will be long term rates of disease a tudy will be the long" term rates of disease a study will be the long" term rates of disease a study will be the long" term rates of disease a tudy will be the long" term rates of disease a tudy will be the long" term rates of disease a tudy will be the long" term rates of disease a tudy will be the long" term rates of disease a tudy will be the l | 119 | The incidence [*] of meningococcal | disease | in England [°] 2 and the United |
| 122IntroductionInvasive meningococcal diseasedisease diseaseis a life threatening condition, ^ than people in employment. were history of preceding induded a history of preceding induded a history of preceding induded a history of preceding in sub-Saharan124with higher* risk of meningococcal of risk factors for meningococcal of risk factors for meningococcal may predispose to meningococcal to risk* factors for meningococcal to risk* factors for meningococcal diseasedisease disease or infuenza. The precise* Discussion129may predispose to meningococcal may predispose to meningococcal to risk* factors for meningococcal diseasedisease disease disease. Further, findings concerning* in adolescents* and a specific in adolescent* and elinical **Outcome measures* Primary and economic productivity of* case mix, *and lis | 120 | in the model, with meningococcal [®] | disease | high season defined as 70 or |
| were at higher risk of meningococcal with higher risk of meningococcal risk factors for meningococcal study^a of epidemic meningococcal of risk factors for meningococcal of risk factors for meningococcal of risk factors for meningococcal to risk factors for meningococcal to risk factors for meningococcal study^a of epidemic meningococcal disease to risk factors for meningococcal to risk factors for meningococcal study^a of risk factors for meningococcal to risk factors for meningococcal study of risk factors for meningococcal study of risk factors for meningococcal to risk factors for meningococcal study of risk factors for meningococcal two winter peaks of meningococcal two winter peaks of meningococcal statutory notifications of meningococcal sease. two winter peaks of meningococcal statutory notifications of meningococcal sease. two winter peaks of meningococcal sease. two winter peaks of meningococcal sease. two winter peaks of meningococcal sease. presentation of recurrent or de novo disease. nadolescents differ^a from those Bigible^a subjects were in teenagers aged 15-19. Of "Experimental and clinical "Ouccome measures^a Primary and economic productivity of^a case mix, "and list size have between^a groups in our study." despite evidence that prevention, family experience of is independent^a of genes risk^a during the adolescent peak canot^a be inferred from the data was sought at the reference "Abuty will be the long^a term rates of disease. of the study will be the long^a term rates of disease. anxiety^a but a substantial effect on disease. | 121 | cohort study of meningococcal | disease | in adolescence to examine |
| 124with higher* risk of meningococcaldiseasewere history of preceding125risk factors for meningococcaldiseaseinduded a history of preceding126study* of epidemic meningococcaldiseaseinduded a history of preceding127for high season of meningococcaldisease"Discussion Risk factors for128of risk factors for meningococcaldisease"Discussion Risk factors for129may predispose to meningococcaldisease"Discussion Risk factors for130to risk* of exposure to meningococcaldisease"Lisellely that our cases were131study of risk factors for meningococcaldiseasein adolescents* and a specific133two winter peaks of meningococcaldiseasein teenagers aged 15-19. Of134statutory notifications of meningococcaldisease"Outcome measures* Primary136presentation of recurrent or de novodisease"Outcome measures* Primary137particularly* given the incidence of disease"Outcome measures* Primary138each practice, including prevalence of disease"Outcome measures* Primary139no differences existed in duration of disease"adependent* of genes140of mannose-binding* lectin and risk of diseaserisk* during the adolescent peak141interest in alternative* types of disease"Results at recruitment form142for a genetic model in which severity of disease"Results at recruitment form143specific epidemiological investigation of disease"Resu | 122 | Introduction Invasive meningocoocal | disease | is a life threatening condition, ^ |
| 125risk factors for meningococcal study ⁶ of epidemic meningococcal to risk factors for meningococcal of risk factors for meningococcal to risk ⁶ of exposure to meningococcal to risk ⁶ of exposure to meningococcal disease to risk ⁶ of exposure to meningococcal disease to risk factors for meningococcal disease to risk ⁶ of exposure to meningococcal disease to risk factors for meningococcal disease tatutory notifications of meningococcal disease to nisk factors for meningococcal di | 123 | were at higher risk of meningococcal [®] | disease | than people in employment. |
| 126study ^a of epidemicmeningococcaldiseasein sub-SaharanAfrica127for high season of meningococcaldiseaseor infuenza. The precise ^a 128of risk factors for meningococcaldiseaseDiscussionRisk factors for129may predispose to meningococcaldisease* DiscussionRisk factors for130to risk ^a of exposure to meningococcaldisease. Further, findings concerning ^a 131study of risk factors for meningococcaldisease. Further, findings concerning ^a 132Risk factors for meningococcaldisease. Bigible ^a subjects were133two winter peaks of meningococcaldisease. Bigible ^a subjects were134statutory notifications of meningococcaldisease. Bigible ^a subjects were136on risk factors for neurodegenerativedisease. Bigible ^a subjects were136presentation of recurrent or de novo disease. "Outcome measures" Primary137particularly ^a given the incidence of disease. "Outcome measures" Primary138each practice, including prevalence of disease. and economic productivity of140of mannose-binding ^a lectin and risk of disease. disease143specific epidemiological investigation of disease. fisedeneett ^a of genes144technique for differing severities of disease. Results at recruitment from145or both). Laboratory confirmation of disease. Results at recruitment from146of the study will be long term rates of d | 124 | with higher ^a risk of meningococcal | disease | were history of preceding |
| 127for high season of meningcocceal of risk factors for meningcocceal may predispose to meningcocceal to risk* of exposure to meningcocceal to risk* of exposure to meningcocceal diseaseor infuenza. The precise* "Discussion Risk factors for t is likely that our cases were . Further, findings concerning* in adolescents* and a specific in adolescents differ* from those131study of risk factors for meningcocceal diseasedisease disease. Further, findings concerning* in adolescents* and a specific in adolescents differ* from those133two winter peaks of meningcocceal diseasedisease disease. Further, findings concerning* in adolescents differ* from those134statutory notifications of meningcocceal diseasedisease disease. Further, findings concerning* in adolescents* and a specific in teenagers aged 15-19. Of * Deperimental and clinical * "Outcome measures* Primary and economic productivity of* case mix, * and list size have between* groups in our study. * despite evidence that prevention, family experience of is independent* of genes risk* during the adolescent peak cannot* be inferred from the data was sought at the reference * Results at recruitment from (APACHE II (acute physiology, The size (about 100 000 specific measures. *12 *13 **14 | 125 | risk factors for meningococcal | disease* | included a history of preceding |
| 128of risk factors for meningcoccealdisease disease* DiscussionRisk factors for t is likely that our cases were130to risk* of exposure to meningcoccealdisease. Further, findings concerning*131study of risk factors for meningcoccealdisease. Further, findings concerning*132Risk factors for meningcoccealdisease. Further, findings concerning*133two winter peaks of meningcoccealdisease. Further, findings concerning*134statutory notifications of meningcoccealdisease. Eligible*subjects were135on risk factors for neurodegenerativedisease. * Durcome measures*Primary136presentation of recurrent or de novo disease.* *Outcome measures*Primary137particularly*given the incidence of disease. * Outcome measures*Primary138each practice, including prevalence of disease. * Courcome measures*Primary139No differences existed in duration of diseaseof mannose-binding*lectin and risk of diseaseexidence that140of mannose-binding*lectin and risk of diseaserisk* during the adolescent peakcanot*be inferred from the data143specific epiderriologicalinvestigation of diseaserisk* during the adolescent peakcanot*be inferred from the data144technique for differing severities of disease* Results at recruitment from(APACHE II (acute physiology,144study will be the long* term rates of disease* Results at recr | 126 | study ^a of epidemic meningococcal | disease | in sub-Saharan Africa |
| 129may predispose to meningcocceal to risk* of exposure to meningcocceal study of risk factors for meningcocceal Risk factors for meningcocceal two winter peaks of meningcocceal statutory notifications of meningcocceal statutory notifications of meningcocceal diseaset is likely that our cases were sease in adolescents* and a specific in adolescents differ* from those133two winter peaks of meningcocceal statutory notifications of meningcocceal on risk factors for neurodegenerative particularly* given the incidence of disease each practice, including prevalence of disease No differences existed in duration of disease to fir a genetic model in which severity of disease specific epidemiological investigation of disease tate or both). Laboratory confirmation of* disease specific epidemiological investigation of* disease specific epidemiological investigation of* disease technique for differing severities of disease tate study will be long term rates of disease study will be the long* term rates of disease. A study will be the long* term rates of disease tate study will be the long* term rates of disease. A anxiety* but a substantial effect on disease specific measures. *12 *13 **14 | 127 | for high season of meningococcal | disease | or influenza. The precise* |
| 130to risk* of exposureto meningcococaldisease).Further, findingsconcerning*131study of risk factorsfor meningcococaldiseasein adolescents*and a specific132Fisk factorsfor meningcococaldiseasein adolescents*and a specific133two winter peaksof meningcococaldiseasein adolescents*and a specific134statutorynotificationsof meningcococaldiseasein teenagersaged 15-19.Of135on risk factorsfor neurodegenerativediseasein teenagersaged 15-19.Of136presentationof recurrent or de novodisease"Outcomemeasures*Primary137particularly*given the incidenceof disease"Outcomemeasures*Primary138each practice, includingprevalenceof diseasebetween*groups in our study."140of mannose-binding*lectinand risk of diseaseprevention, family experience ofis independent*of genes142for a geneticmodel in which severityof diseaserisk* duringthe adolescentpeak144techniquefor differingseveritiesof disease"Resultsat recruitmentform144techniquefor differingseverity*of disease"Resultsat recruitmentform145or both)Laboratoryconfirmationof disease"Resultsat recruitmentfor | 128 | of risk factors for meningococcal | disease | ^a Discussion Risk factors for |
| 131study of risk factors for meningcooccaldiseasein adolescents*and a specific132Risk factors for meningcooccaldiseasein adolescents*and a specific133two winter peaks of meningcooccaldiseasein adolescents*differ*from those134statutory notifications of meningcooccaldiseasein teenagers aged 15-19. Of135on risk factors for neurodegenerativedisease* Experimentaland clinical136presentation of recurrent or de novodisease* Outcome measures*Primary137particularly*given the incidence of diseaseand economic productivity of*138each practice, including prevalence of diseasecase mix, * and list size have139No differences existed in duration of diseasebetween*groups in our study. *140of mannose-binding*lectin and risk of diseaserisk* during the adolescent peak141interest in alternative*types of diseaserisk* during the adolescent peak143specific epidemiological investigation of diseaserisk* during the adolescent peak144technique for differing severities of disease* Results at recruitment form147by age, sex, HADS score, severity* of disease* Results at recruitment form148study will be the long* term rates of disease* Results at recruitment form149anxiety* but a substantial effect on diseaseThe size (about 100 000149anxiety* but a substantial effect on diseasethe size <td>129</td> <td>may predispose to meningococcal*</td> <td>disease.</td> <td>t is likely that our cases were</td> | 129 | may predispose to meningococcal* | disease. | t is likely that our cases were |
| 132Risk factors for meningococcaldiseasein adolescentsdiffer^n from those133two winter peaks of meningococcaldiseasein teenagers aged 15-19. Of134statutory notifications of meningococcaldiseasein teenagers aged 15-19. Of135on risk factors for neurodegenerativedisease* Dutcome measures* Primary136presentation of recurrent or de novo disease.* Outcome measures* Primary137particularly^ given the incidence of disease.and economic productivity of*138each practice, including prevalence of disease.case mix, * and list size have139No differences existed in duration of disease.between* groups in our study. *140of mannose-binding* lectin and risk of disease.prevention, family experience of142for a genetic model in which severity of disease.risk* during the adolescent peak144technique for differing severities of disease.* Results at recruitment form145or both). Laboratory confirmation of* disease.* Results at recruitment form147by age, sex, HADS score, severity* of disease.* Results at recruitment form148study will be the long* term rates of disease.* Results at recruitment form149anxiety* but a substantial effect on disease.* Results.* 12 *13 **14 | 130 | to risk [®] of exposure to meningococcal | disease) | . Further, findings concerning* |
| 133two winter peaks of meningococcaldisease.Eligible* subjects were134statutory notifications of meningococcaldisease*in teenagers aged 15-19. Of135on risk factors for neurodegenerativedisease** Experimentaland clinical136presentation of recurrent or de novo disease.* "Outcome measures*" Primary137particularly*given the incidence of disease.* "Outcome measures*" Primary138each practice, including prevalence of disease.case mix, * and list size have139No differences existed in duration of disease.case mix, * and list size have140of mannose-binding*lectin and risk of disease.teween*141interest in alternative*types of disease.revention, family experience of142for a genetic model in which severity of disease.risk* during the adolescent peak144technique for differing severities of disease.* Results at recruitment form145or both)Laboratory confirmation of* disease.* Results at recruitment form147by age, sex, HADS score, severity* of disease.* Results at recruitment form148study will be the long* term rates of disease.* Results at recruitment form149anxiety* but a substantial effect on disease.* Results.* 12 *13 **14 | 131 | study of risk factors for meningococcal | disease | in adolescents" and a specific |
| 134statutorynotificationsof meningococcaldiseasein teenagersaged 15-19.Of135on risk factorsfor neurodegenerativedisease* Experimentaland clinical136presentationof recurrentor de novodisease.* *Outcomemeasures*Primary137particularlygiventhe incidenceof disease.* *Outcomemeasures*Primary138eachpractice,includingprevalenceof disease,casemix, * and list sizehave139Nodifferencesexistedin durationof disease,casemix, * and list sizehave140of mannose-binding*lectinand risk of disease,despiteevidencethat141interestin alternative*typesof disease,is independent*of genes142for a geneticmodelin whichseverityof disease,risk* duringthe adolescentpeak144techniquefor differingseveritiesof disease,risk* duringthe adolescentpeak145or both)Laboratoryconfirmationof disease,* Resultsat recruitmentform146of the study will be long termrates of disease,* Resultsat recruitmentform147by age, sex, HADSscore, severity*of disease,* Resultsat recruitmentform148study will be the long* termrates of disease,< | 132 | Risk factors for meningococcal | disease | in adolescents differ ^a from those |
| 134statutorynotificationsof meningococcaldiseasein teenagersaged 15-19.Of135on risk factorsfor neurodegenerativedisease* Experimentaland clinical136presentationof recurrentor de novodisease.* Outcomemeasures*Primary137particularly*giventhe incidenceof disease.* *Outcomemeasures*Primary138eachpractice,includingprevalenceof disease.and economicproductivityof*138eachpractice,includingprevalenceof disease.casemix, * and list sizehave139Nodifferencesexistedin durationof disease.between*groups in our study. *140of mannose-binding*lectinand risk of diseaseprevention, familyexperienceof141interest in alternative*typesof diseaseis independent*of genes143specificepiderniologicalinvestigationof diseaserisk* duringthe adolescentpeak144techniquefor differingseveritiesof disease"Resultsat recruitmentform145or both)Laboratoryconfirmationof disease."Resultsat recruitmentform146of the study will be long termrates of disease."Resultsat recruitmentform147by age, sex, HADSscore, severity*of disease.< | 133 | two winter peaks of meningcooccal | disease. | Bigible^ subjects were |
| 135on risk factors for neurodegenerativedisease.* Experimentaland clinical136presentationof recurrentor de novodisease.* *Outcomemeasures*Primary137particularly*given the incidenceof diseaseand economicproductivityof*138each practice, includingprevalenceof diseaseand economicproductivityof*139No differencesexisted in durationof diseasebetween*groups in our study. *140of mannose-binding*lectinand risk of diseasedespiteevidencethat141interest in alternative*typesof diseaseprevention, family experienceof142for a geneticmodel in which severityof diseaseis independent*of genes143specificepidemiologicalinvestigationof diseaserisk* during the adolescent peak144techniquefor differingseveritiesof diseaseansouth be inferred from the data145or both)Laboratoryconfirmationof*disease* Results at recruitment146of the study will be long term ratesof disease* Results at recruitmentform147by age, sex, HADSscore, severity*of diseaseThe size (about 100 000148anxiety*but a substantialeffect on diseasespecificmeasures.*12 **14 | 134 | · · · | disease* | • , |
| 136presentationof recurrentor de novodisease.^*Outcomemeasures*Primary137particularly^giventhe incidenceof diseaseand economicproductivityof'138eachpractice,includingprevalenceof disease,casemix, ^andlistsizehave139Nodifferencesexistedindurationof disease,casemix, ^andlistsizehave140of mannose-binding^lectinand risk of diseasebetween^groupsin our study. ^141interestin alternative^typesof diseaseprevention,familyexperienceof142for a geneticmodelin whichseverityof diseaseisindependent^of genes143specificepidemiologicalinvestigationof diseaserisk^duringthe adolescentpeak144techniquefor differingseveritiesof diseasecannot^be inferredformthe data145or both)Laboratoryconfirmationof disease"Resultsat recruitmentform147by age, sex, HADSscore, severity^of disease."Resultsat recruitmentform148studywill be the long^termratesof disease.The size(about100000149anxiety^but a substantialeffecton diseasespecific< | 135 | · · | disease. | |
| 137 particularlyⁿ given the incidence of disease and economic productivity of 138 each practice, including prevalence of disease, case mix, ⁿ and list size have 139 No differences existed in duration of disease betweenⁿ groups in our study. ⁿ 140 of mannose-bindingⁿ lectin and risk of disease despite evidence that 141 interest in alternativeⁿ types of disease is independentⁿ of genes 143 specific epidemiological investigation of disease of disease cannotⁿ be inferred from the data 144 technique for differing severities of disease of disease is sought at the reference 145 or both). Laboratory confirmation of disease. ⁿ Results at recruitment from 147 by age, sex, HADS score, severityⁿ of disease. ⁿ Results at recruitment from 148 study will be the longⁿ term rates of disease. The size (about 100 000 149 anxietyⁿ but a substantial effect on disease specific measures. ⁿ12 ⁿ13 ⁿ14 | 136 | | disease. | • |
| 138eachpractice,includingprevalenceof disease,casemix, ^ andlist sizehave139Nodifferencesexistedindurationof diseasebetween^groupsinour study. ^140ofmannose-binding^lectinandriskof diseasedespiteevidencethat141interestin alternative^typesof diseaseprevention,familyexperienceof142for a geneticmodelin whichseverityof diseaseisindependent^of genes143specificepidemiologicalinvestigationof diseaserisk^duringthe adolescentpeak144techniquefor differingseveritiesof diseasecannot*be inferredfromthe data145or both)Laboratoryconfirmationof*diseaseass soughtat the reference146of the study will be long termratesof disease(APACHEII (acutephysiology,148study will be the long* termratesof diseaseThe size (about100000149anxiety*but a substantialeffectondiseasespecificmeasures.*12*13*14 | 137 | • | disease | , |
| 139No differencesexistedin durationof diseasebetween*groupsin our study.*140of mannose-binding*lectinand riskof diseasebetween*groupsin our study.*141interestin alternative*typesof diseasedespiteevidencethat142for a geneticmodelin whichseverityof diseaseis independent*of genes143specificepidemiologicalinvestigationof diseaserisk* duringthe adolescentpeak144techniquefor differingseveritiesof diseasecannot*be inferredfromthe data145or both)Laboratoryconfirmationof* disease*Resultsat recruitmentfrom147by age, sex,HADSscore,severity*of disease.*Resultsat recruitmentfrom148studywill be the long* termratesof disease.The size(APACHEII (acutephysiology,149anxiety*but a substantialeffectondiseasespecificmeasures.*12*13*14 | 138 | · · · | disease. | · · · |
| 140of mannose-binding*lectinand riskof diseasedespiteevidencethat141interestin alternative*typesof diseaseprevention,familyexperienceof142for a geneticmodelin whichseverityof diseaseis independent*of genes143specificepidemiologicalinvestigationof diseaserisk* duringthe adolescentpeak144techniquefor differingseveritiesof diseasecannot*be inferredfromthe data145or both)Laboratoryconfirmationof*diseasewassoughtat the reference146of the study will be long termratesof disease*Resultsat recruitmentfrom147by age, sex,HADSscore,severity*of disease.(APACHEII (acutephysiology,148study will be the long* termratesof disease.The size (about100000149anxiety*but a substantialeffectondiseasespecificmeasures.*12*14 | | | | • |
| 141interest in alternative* types of diseaseprevention, family experience of142for a genetic model in which severity of diseaseis independent* of genes143specific epidemiological investigation of diseaserisk* during the adolescent peak144technique for differing severities of diseasecannot* be inferred from the data145or both)Laboratoryconfirmation146of the study will be long term ratesof disease* Results147by age, sex, HADSscore, severity* of disease(APACHE II (acute physiology,148study will be the long* term ratesof diseaseThe size (about 100 000149anxiety* but a substantialeffect on diseasespecific measures.*12 **14 | 140 | of mannose-binding^lectinand_risk_of | disease | • • • • |
| 142for a genetic model in which severity of diseaseis independent*of genes143specific epidemiological investigation of diseaserisk* during the adolescent peak144technique for differing severities of diseasecannot* be inferred from the data145or both)Laboratoryconfirmationof disease146of the study will be long term rates of disease* Results at recruitment from147by age, sex, HADS score, severity* of disease(APACHE II (acute physiology,148study will be the long* term rates of disease.The size (about 100 000149anxiety* but a substantial effect on diseasespecific measures.*12 **13 **14 | | • | | • |
| specific epidemiological investigation of disease risk^a during the adolescent peak technique for differing severities of disease cannot^a be inferred from the data or both) . Laboratory confirmation of^a disease was sought at the reference of the study will be long term rates of disease. ^a Results at recruitment from by age, sex, HADS score, severity^a of disease. The size (about 100 000 study will be the long^a term rates of disease. The size (about 100 000 anxiety^a but a substantial effect on disease specific measures. ^a12 ^a13 ^a14 | 142 | 2 | | |
| 144 technique for differing severities of disease cannot [*] be inferred from the data 145 or both) . Laboratory confirmation of disease was sought at the reference 146 of the study will be long term rates of disease. * Results at recruitment from 147 by age, sex, HADS score, severity [*] of disease (APACHE II (acute physiology, 148 study will be the long [*] term rates of disease. The size (about 100 000 149 anxiety [*] but a substantial effect on disease specific measures. *12 *13 **14 | | | | |
| or both) Laboratory confirmation of disease was sought at the reference of the study will be long term rates of disease. "Results at recruitment from by age, sex, HADS score, severity" of disease (APACHE II (acute physiology, study will be the long" term rates of disease. The size (about 100 000 anxiety" but a substantial effect on disease specific measures. "12 "13 "14 | | | | |
| 146 of the study will be long term rates of disease. ^ Results at recruitment from 147 by age, sex, HADS score, severity^ of disease (APACHE II(acute physiology, 148 study will be the long^ term rates of disease. The size (about 100 000 149 anxiety^ but a substantial effect on disease specific measures. ^12 ^13 ^^14 | | | | |
| 147 by age, sex, HADS score, severity^ of disease (APACHE II(acute physiology, 148 study will be the long^ term rates of disease. The size (about 100 000 149 anxiety^ but a substantial effect on disease specific measures. ^12 ^13 ^^14 | | | | - |
| 148 study will be the long ^a term rates of disease. The size (about 100 000 149 anxiety ^a but a substantial effect on disease specific measures. ^a 12 ^a 13 ^a 14 | | | | |
| 149 anxiety ^a but a substantial effect on disease specific measures. ¹² ¹³ ¹⁴ | | | | |
| | | | | |
| The management of entering provided discuss is variable, contentions, allo | | | | |
| | | | | |

SEZIONE 1

| N | Concordance |
|------------|--|
| 151 | treatment and management of pilonidal disease^ is required. Evidence is still |
| 152 | study of invasive pneumococcal disease^ by Nuorti et al.^49 However, |
| 153 | 2 diabetes mellitus is a progressive disease, which can' lead to considerable |
| 154 | vascular accident, chronic pulmonary disease, urinary incontinence,^ |
| 155 | severe chronic cardiac or pulmonary disease (New York Heart' Association |
| 156 | cancer, chronic obstructive pulmonary disease, peripheral vascular disease. |
| 157 | or chronic obstructive pulmonary disease, which do not require treatment |
| 158 | vascular accident, chronic pulmonary disease, which do not require treatment |
| 150 | (ADS),^ the gastro-oesophageal reflux disease-health related quality^ of life |
| 160 | in patients with^ end stage renal disease treated by haemodialysis was |
| 161 | intracerebral haemorrhage, renal disease, liver disease,^ dementia or |
| 162 | led to declining coverage and resultant disease outbreaks in^ the United |
| 163 | disorders,^ liver disease, rheumatic disease, medical school affiliation,^ other |
| 164 | tendinopathy, Osgood-Schlatter ^A disease, or other defined pathological |
| | |
| 165 | severely affected patients. More severe disease presumably indicates a greater |
| 166 | shock^ wave therapy for shoulder disease.^34 ^35 ^36 Thus, our results^ |
| 167 | to severe discomfort or pain, skin disease associated^ with cold |
| 168 | histological high grade squamous^ disease than did manually read |
| 169 170 | is inappropriate?for^ example, terminal disease or patient choice?a patient^ may |
| | with the first character indicating [^] the disease area and later characters |
| 171 | tend to appear late in the course of the disease, and early surgery remains the |
| 172 | Awareness [^] and understanding of the disease Perceived seriousness of the [^] |
| 173 | of severity they associated with the disease. All newly diagnosed patients |
| 174 | to several perceptions and factors: the disease having been discovered "at an |
| 175 | type 2 diabetes/ Perceptions^ of the disease (including perceived |
| 176 | and mortality associated with the disease, our systematic review found few |
| 177 | disease Ideas about the causes of the disease /After results^ of the oral |
| 178 | disease Perceived seriousness of the disease Ideas about the causes of the |
| 179 | the potential consequences ⁴ of the disease to justify lifestyle change, |
| 180 | those ventually diagnosed with the disease. A This approximate participants talked |
| 181 | high readings to progression of the disease. ^A This association between self |
| 182 | being^ a good thing?enabling the disease to be detected at an^ early, |
| 183 | about^ their plans to control the disease; in some cases a diet-only^ |
| 184 | expressed in their plans to control their disease^ seemed to be related to several |
| 185 | and their experiences of managing their disease.^20 ^21 ^22 ^23 ^ 24 ^25 |
| 186 | younger populations and therefore this disease has an economic impact. To |
| 187 | the alleles assumed to predispose [^] to disease. [^] 19 [^] 20 [^] 21 Another prediction of |
| 188 | pulmonary^ disease, peripheral vascular disease, psychiatric disorders,^ liver |
| 189 | disease, peripheral vascular disease, coronary heart disease, heart |
| 190 | disease, peripheral vascular disease,^ coronary heart disease, heart |
| 191 | techniques. Heart^ failure, vascular disease, and diabetes were defined by |
| 192 | disease, peripheral vascular disease, coronary heart disease, heart |
| 193 | for more advanced diabetes or vascular disease (or both).^w6 ^ Thus, treatment |
| 194 | lectin was not associated with disease. Immunisation [^] was protective, |
| | |
| | |

Figure 2. Concordance of *illness* in the corpus from the BMJ (109,822), sorted by the first word to the left.

N Concordance (compared with those diagnosed after^ illness)^7 ^8 and a lack of understanding 1 or had a psychotic illness^ or an illness with a prognosis of less than one 2 3 We reduced the time[^] between illness and interview and used memory 4 in a patient with a clinically compatible illness.[^] Consultants in communicable 5 rehabilitation requirements[^] after critical illness has advanced since we designed recovery and[^] rehabilitation after critical illness is also required. Finally,[^] the role 6 7 23 May 2005), as apparent flu-like illness or hangover^ may in fact be the 8 to hospital for an acute medical illness.^4 ^ ^13 These studies agreed 9 to intensive care or symptoms of illness.[^] In the univariate matched 10 subgroups,^ which included severity of illness, chronic comorbidity, intensive^ undertaken for APACHE II severity of illness, ^ APACHE II comorbidity, 11 12 may be related to the lower severity of illness^ in the patients in the second may be reduced by the high severity of illness of patients treated^ in UK 13 14 the month before the case patient's illness, which was has easier to do as their 15 data from the participants on preceding illness:^ symptoms of prodromal illness disease were history of preceding illness, ^ intimate kissing, being a 16 to report being unwell^ with a preceding illness in the fortnight before admission 17 (in 15-19 year olds).^ *Role of preceding illness* Independent biological risk 18 19 included a history of preceding illness and preterm birth. A^ preceding of preceding illness. Preceding illness occurred in 53% of cases^ and 20 symptom based definition[^] of preceding illness. Preceding illness occurred in 21 The precise[^] aetiology of this preceding illness is unclear and may be a[^] 22 23 illness and preterm birth. A^ preceding illness has previously been identified as 24 illness (in cases) and preceding illness^ (in cases and controls?for 25 illness: ^ symptoms of prodromal illness (in cases) and preceding illness ^ because of preceding or prodromal^illness. Religious observance has been 26 27 bias arises from preceding or prodromal illness, as a reduction^ in risk behaviours or lactating, or had a psychotic illness^ or an illness with a prognosis of 28 29 for important aspects of a patient?s illness^ such as delirium and cognitive 30 speaking,^ altered mental state, severe illness, urgent need for cannulation),^ indicated that' behaviour change due to illness was not a significant source' of 31 sustaining health rather^ than treating illness."^13 ^ We included trials reporting 32 33 creatinine >221 mmol/l; unstable illness;^ active ischaemia *Agents

Appendix 2 (a)

The following is a selection of text pieces - in the corpus from the BMJ online (95,288 words) - where *disease* is associated with specific pathological conditions.

Names of pathological conditions and occurrences of *disease* have been highlighted in bold type.

1. Unsurprisingly, the striking feature is the numbers of infectious and parasitic **diseases**, which accounted for nearly a fifth of deaths. Many of these were in the young, with over 90% of deaths from diarrhoea and **dysentery**, **measles**, and **whooping cough** being among those aged under 5. (BMJ 2009;339:b3454, doi: 10.1136/bmj.b3454)

2. Self reports of major chronic diseases (such as cancer, diabetes, coronary heart disease, stroke, Parkinson's disease, and multiple sclerosis) were confirmed through various methods, including review of medical records and pathology reports, telephone interview, and supplementary questionnaires to participants.

(BMJ 2006;332:875-884 (15 April), doi:10.1136/bmj.38771.583796.7C)

3. Our exclusion criteria were a history of major chronic **diseases** at study baseline in 1976, including **cancer**, **diabetes**, **myocardial infarction**, coronary artery bypass graft surgery, **stroke**, **kidney failure**, **chronic obstructive pulmonary disease**, **Parkinson's disease**.

(BMJ 2009;339:b3796, doi: 10.1136/bmj.b3796)

4. If **pneumonia** was suspected the participant was assessed at the affiliated hospital; the medical staff at the nursing homes also informed the study coordinator of the occurrence of **pneumonia** or any other **disease**. (BMJ 2010;340:c1004, doi: 10.1136/bmj.c1004)

5. Interpreting the relevance of associations between low body mass index and chronic **disease** such as **cirrhosis** is difficult, as analyses may not adequately compensate for the likelihood that early liver disease may affect body mass index before the first hospital admission or death occurs -for example, by reducing appetite or by causing malabsorption. (BMJ 2010;340:c912, doi: 10.1136/bmj.c912)

6. The challenge of treating patients with both active **psoriasis** and active **psoriatic arthritis** is to optimise the treatment of both **disease** manifestations to give the best overall outcome.

(BMJ 2010;340:c147, doi: 10.1136/bmj.c147)

7. The cutaneous symptoms of **psoriatic arthritis** usually appear a decade or more before the joint symptoms, enthesitis, and dactylitis. The **disease** affects men and women equally and has a worldwide distribution. (BMJ 2009;339:b2433, doi: 10.1136/bmj.b2433)

8. 20. Box 1: Diseases included in disease count

* Hypertension

* Ischaemic heart disease

* Cerebrovascular disease

* Peripheral vascular disease

* Heart failure

* Atrial flutter or fibrillation

* Arthritis (osteoarthritis or cervical or lumbar spondylosis or rheumatoid arthritis or other arthritis or non-specified arthritis)

* Osteoporosis

* Chronic obstructive pulmonary disease or asthma

* Diabetes

* Hypothyroidism or hyperthyroidism

* Cancer diagnosed within past five years (excluding non-melanoma skin cancer)

* Eye disease (cataract or age related macular degeneration or glaucoma or diabetic eye disease or registered blind or partially sighted)

- * Dementia
- * Parkinson's disease
- * Renal impairment

(BMJ Apr 2008; 336: 754 - 757; doi:10.1136/bmj.39489.590671.25)

Appendix 2 (b)

The following is a selection of text pieces - in the corpus from the BMJ (95,288 words) - where *illness* occurs. In all extracts selected *illness* does not refer to any specific pathological condition, except in extracts 6 and 7.

Names of pathological conditions and occurrences of *illness* have been highlighted in bold type.

1. Compared with the reference group (women with a body mass index of 22.5 to <25), both the women who had a lower body mass index and those with a higher body mass index had a significantly greater relative risk of cirrhosis. Among women with body mass index below 22.5, we cannot exclude the possibility that previous **illness** may have contributed to weight loss, and for this reason we focused our analyses on women with a body mass index of 22.5 or above. Among women who had a body mass index of 22.5 or above, little evidence existed to suggest non-linearity in the relation between body mass index and the relative risk of cirrhosis related hospital admission or death (test for non-linearity, P=0.2). (BMJ 2010;340:c912, doi: 10.1136/bmj.c912)

2. We invited mothers who gave birth at the maternity wards of the national hospital and a local health centre (Bandim Health Centre) to participate in the study when their child was to receive BCG vaccination after delivery. Furthermore, we invited mothers who delivered at home to participate when they came for BCG vaccination at two of the three health centres in the study area (Bandim and Belem). We did not include the third health centre (Cuntum) for logistical reasons. The inclusion criteria were weight at least 2500 g at presentation and no signs of overt **illness** or malformations. Infants who died in the maternity wards, before the vaccination team arrived in the morning, could not be assessed for enrolment. This was also the case for those born at home who died before their mother brought them for vaccination.

(BMJ Jun 2008; 336: 1416 - 1420; doi:10.1136/bmj.39542.509444.AE)

3. To examine the relation between hospital orthopaedic specialisation and postoperative surgical outcomes we carried out a retrospective cohort study of US Medicare beneficiaries (all Americans aged 65 and older) who underwent total hip replacement or total knee replacement. We developed a measure of a hospital's orthopaedic specialisation and then compared the demographics, socioeconomic status, and prevalence of comorbid **illness** among patients who received total hip replacement or total knee replacement in hospitals with lower and higher degrees of orthopaedic specialisation. We then assessed the structural characteristics of less and more specialised hospitals.

(BMJ 2010;340:c165, doi: 10.1136/bmj.c165)

4. In contrast, the effects of illness or drug use on sexual function in women are poorly understood. Sexual problems, including low desire, vaginal dryness, difficulties with orgasm, and pain with intercourse are prevalent among sexually active older women,1 are associated with decreased sexual satisfaction, but typically do not render a woman physically incapable of sexual intercourse. Women's sexual interest or motivation may be more resilient to **illness** or sexual problems than men's, may be more contextually dependent on the partner or situational factors, or, as seen in younger populations, older women may have less agency over their sexual activity than men.

(BMJ 2010;340:c810, doi: 10.1136/bmj.c810)

5. To examine the relation between hospital orthopaedic specialisation and postoperative surgical outcomes we carried out a retrospective cohort study of US Medicare beneficiaries (all Americans aged 65 and older) who underwent total hip replacement or total knee replacement. We developed a measure of a hospital's orthopaedic specialisation and then compared the demographics, socioeconomic status, and prevalence of comorbid **illness** among patients who received total hip replacement or total knee replacement in hospitals with lower and higher degrees of orthopaedic specialisation.

(BMJ 2010;340:c165, doi: 10.1136/bmj.c165)

6. Pregnant women are at increased risk of influenza and its complications. The effects of influenza during pregnancy have been noted in previous pandemics, particularly the increased mortality in pregnant women compared with the general population. The 2009 influenza A/H1N1 pandemic was the first influenza pandemic to occur in the era of modern obstetric and intensive care management, and pregnancy is a risk factor for critical **illness** due to 2009 **H1N1 influenza**. (...)

Of 59 women with available data, 25 (39%) had a body mass index of more than 30 and 13 (20%) an index or more than 35. Overall, 36 women (56%) had a documented coexisting **illness**, of whom 21 (33% of total) had **asthma**. (BMJ 2010;340:c1279, doi: 10.1136/bmj.c1279)

7. The 3818 study hospitals carried out 1 273 081 major joint replacements from 2001-5. Patient characteristics and comorbidity varied as hospital specialisation increased (table 1Go). In particular, more specialised hospitals treated a lower proportion of women and African-Americans (P<0.001 for each) and those with fewer comorbid **illnesses**, including **diabetes** and **heart failure**.

(...)

Myalgic encephalitis, is characterised by disabling physical and mental fatigue, lasting for at least six months, without an apparent physical cause. The hallmark of the **illness** is debilitating fatigue, but symptoms like myalgia, disrupted sleep, difficulty with concentration, sore throat, and lymphadenopathy may also be present, albeit more variably. More than two thirds of patients are women. Although the cause is unknown and the **illness** may cover more than one entity, many have suggested that infectious agents have a role. (BMJ 2010;340:c165, doi: 10.1136/bmj.c165)