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DATA DESCRIPTOR

BioASQ-QA: A manually curated corpus for Biomedical Question Answering

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The BioASQ question answering (QA) benchmark dataset contains questions in English, along with golden standard (reference) answers and related material. The dataset has been designed to reflect real information needs of biomedical experts and is therefore more realistic and challenging than most existing datasets. Furthermore, unlike most previous QA benchmarks that contain only exact answers, the BioASQ-QA dataset also includes ideal answers (in effect summaries), which are particularly useful for research on multi-document summarization. The dataset combines structured and unstructured data. The materials linked with each question comprise documents and snippets, which are useful for Information Retrieval and Passage Retrieval experiments, as well as concepts that are useful in concept-to-text Natural Language Generation. Researchers working on paraphrasing and textual entailment can also measure the degree to which their methods improve the performance of biomedical QA systems. Last but not least, the dataset is continuously extended, as the BioASQ challenge is running and new data are generated.

Background & Summary

More than 2 articles are published in biomedical journals every minute, leading to MEDLINE/PubMed¹ currently comprising more than 32 million articles, while the number and size of non-textual biomedical data sources also increases rapidly. As an example, since the outbreak of the COVID-19 pandemic, there has been an explosion of new scientific literature about the disease and the virus that causes it, with about 10,000 new COVID-19 related articles added each month². This wealth of new knowledge plays a central role in the progress achieved in biomedicine and its impact on public health, but it is also overwhelming for the biomedical expert. Ensuring that this knowledge is used for the benefit of the patients in a timely manner is a demanding task.

BioASQ³ (Biomedical Semantic Indexing and Question Answering) pushes research towards highly precise biomedical information access systems through a series of evaluation campaigns, in which systems from teams around the world compete. BioASQ campaigns run annually since 2012, providing data, open-source software and a stable evaluation environment for the participating systems. In the last ten years that the challenge has been running, around 100 different universities and companies, from all continents, have participated in BioASQ, providing a competitive, but also synergetic ecosystem. The fact that the participants of the BioASQ challenges are all working on the same benchmark data, facilitates significantly the exchange and fusion of ideas and eventually accelerates progress in the field. The ultimate goal is to lead biomedical information access systems to the maturity and reliability required by biomedical researchers.

BioASQ comprises two main tasks. In Task A systems are asked to automatically assign Medical Subject Headings (MeSH)⁴ terms to biomedical articles, thus assisting the indexing of biomedical literature. Task B focuses on obtaining precise and comprehensible answers to biomedical research questions. The systems that participate in Task B are given English questions that are written by biomedical experts and reflect real-life information needs. For each question, the systems are required to return relevant articles, snippets of the articles, concepts from designated ontologies, RDF triples from Linked Life Data⁵, an 'exact' answer (e.g., a disease or symptom), and a paragraph-sized summary answer. Hence, this task combines traditional information retrieval, with question answering from text and structured data, as well as multi-document text summarization.

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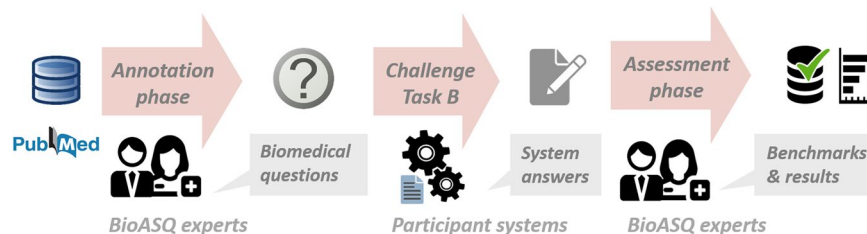


Fig. 1 During the annotation phase of the BioASQ, the experts compose biomedical questions. The participating systems provide answers in the challenge. Finally, in the assessment phase, the experts manually assess the system responses and refine and extend the dataset.

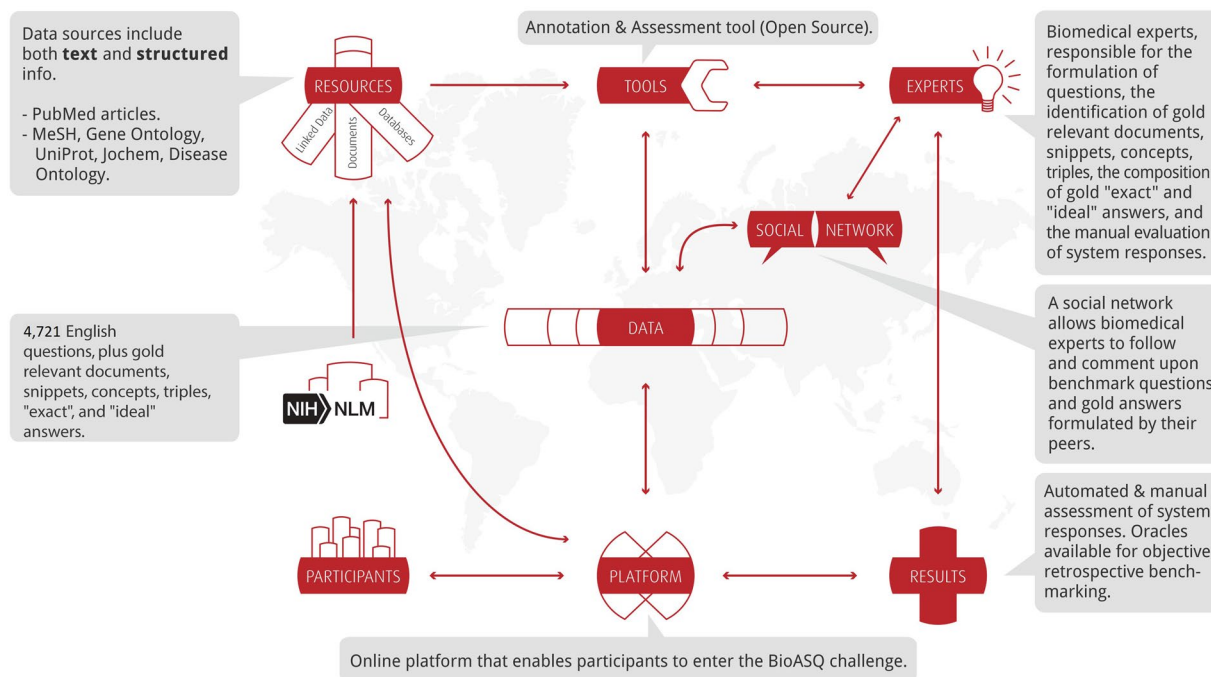


Fig. 2 The BioASQ infrastructure and ecosystem.

One of the main tangible outcomes of BioASQ is its benchmark datasets. The BioASQ-QA dataset that is generated for Task B, contains questions in English, along with golden standard (reference) answers and supporting material. The BioASQ data are more realistic and challenging than most existing datasets for biomedical expert question answering^{6,7}. In order to achieve this, BioASQ employs a team of trained experts, who provide annually a set of around 500 questions from their specialized field of expertise. Figure 1 provides the lifecycle of the BioASQ dataset creation, which is presented in detail in the following sections. Using this process, a set of 4721 questions and answers have been generated so far, constituting a unique resource for the development of QA systems.

Methods

The BioASQ infrastructure and ecosystem. Figure 2 summarises the main components of the BioASQ infrastructure, as well as key stakeholders in the related ecosystem. The BioASQ infrastructure includes tools for annotating data, tools for assessing the results of participating systems, benchmark repositories, evaluation services, etc. The infrastructure allows challenge participants to access training and test data, submit their results and be informed about the performance of their systems, in comparison to other systems. The BioASQ infrastructure is also used by the experts during the creation of the benchmark datasets and helps improve the quality of the data. In the following subsections, the different components of the BioASQ ecosystems are described.

Expert team. As the goal of BioASQ is to reflect real information needs of biomedical experts, their involvement was necessary in the creation of the dataset. The biomedical expert team of BioASQ was first established in 2012, but has changed through the years. Several experts were considered at that time, from a variety of institutions across Europe. The final selection of the experts was based on the need to cover the broad biomedical scientific field, representing as much as possible, medicine, biosciences and bioinformatics. The members of the biomedical team hold positions in universities, hospitals or research institutes in Europe. Their primary research

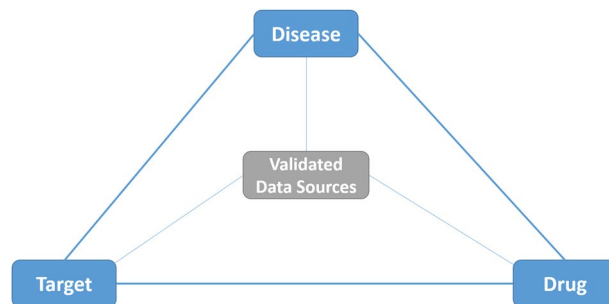


Fig. 3 The drug-target-disease triangle, adopted in *BioASQ*.

interests include: cardiovascular endocrinology, psychiatry, psychophysiology, pharmacology, drug repositioning, cardiac remodeling, cardiovascular pharmacology, computational genomics, pharmacogenomics, comparative genomics, molecular evolution, proteomics, mass spectrometry, protein evolution, clinical information retrieval from electronic health records, and clinical practice guidelines. In total 21 experts have contributed to the creation of the dataset, 7 of whom have been involved most actively. The main job of the biomedical expert team is the creation of the QA benchmark dataset, using an annotation tool provided by BioASQ. With the use of the tool, the experts can set their questions and retrieve relevant documents and snippets from MEDLINE. Additionally, the biomedical expert team assesses the responses of the participating systems. In addition to scoring the systems' answers, during this process the experts have the opportunity to enrich and modify the gold material that they have provided, thus improving the quality of the benchmark dataset.

Regular physical and virtual meetings are organised with the experts. Partly, these meetings aim to train the new members of the team and inform the existing ones about changes that have happened. In particular, the goals of the training sessions are as follows:

- Familiarization with the annotation and assessment tools used during the formulation and assessment of biomedical questions respectively. This step also involves familiarization of the experts with the specific types of questions used in the challenge, i.e. factoid, yes/no, list and summary questions. At the same time, the experts provide feedback and help shaping the BioASQ tools.
- Familiarization with the resources used in BioASQ, both MEDLINE and various structured sources. The aim is to help the experts understand the data provided by these source, in response to different questions they may formulate.
- Resolution of issues that come up during the question composition and assessment tasks. This is a continuous process that extends beyond the training sessions. Continuous support is provided to the experts, while the experts can also interact with each other and provide feedback on the data being created.

Data selection. The QA benchmark is based primarily on documents indexed for *MEDLINE*. In addition, a wide range of biomedical concepts are drawn from ontologies and linked data that describe different facets of the domain. The selected resources follow commonly used *drug-target-disease triangle*, which defines the prime information axes for medical investigations. The main principle is shown in Figure 3.

This “*knowledge-triangle*” supports the conceptual linking of biomedical knowledge databases and related resources. Based on this, systems can address questions, linking natural language questions with relevant ontology concepts. In this context, the following resources have been selected for BioASQ.

Drugs: **Jochem**⁸, the Joint Chemical Dictionary, is a dictionary for the identification of small molecules and drugs in text, combining information from UMLS, MeSH, ChEBI, DrugBank, KEGG, HMDB, and ChemIDplus. Given the variety and the population of the different resources in it, Jochem is currently one of the largest biomedical resources for drugs and chemicals.

Targets: **Gene Ontology (GO)**^{9,10} is currently the most successful case of ontology use in bioinformatics and provides a controlled vocabulary to describe functional aspects of gene products. The ontology covers three domains: cellular component, molecular function, and biological process.

Universal Protein Resource (UniProt)¹¹ provides a comprehensive, high-quality and freely accessible resource of protein sequence and functional information. Its protein knowledge base consists of two sections: SwissProt, which is manually annotated and reviewed, and contains more than 500 thousand sequences, and TrEMBL, which is automatically annotated and is not reviewed, and contains a few million sequences. In BioASQ the SwissProt component of UniProt is used.

Diseases: **Disease Ontology (DO)**¹² contains data associating genes with human diseases, using established disease codes and terminologies. Approximately 8,000 inherited, developmental and acquired human diseases are included in the resource. The DO semantically integrates disease and medical vocabularies through extensive cross-mapping and integration of MeSH, ICD, NCI's thesaurus, SNOMED CT and OMIM disease-specific terms and identifiers.

Document Sources: The main source of biomedical literature is NLM's MEDLINE and is accessible through PubMed and PubMed Central. PubMed, indexes over 34 million citations, while PubMed Central (PMC) provides free access to approximately 8.5 million full-text biomedical and life-science articles.

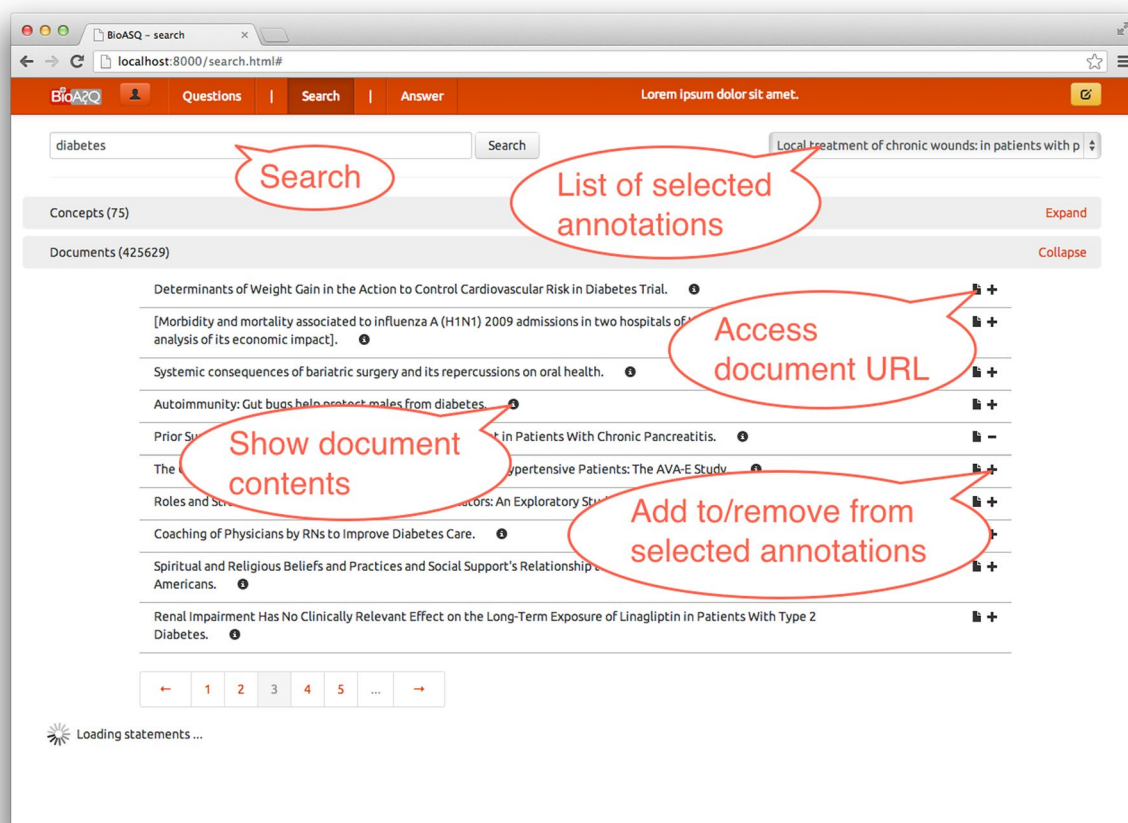


Fig. 5 Screenshot of the annotation tool's search and data selection screen with the section for *document results* expanded.

with a one-paragraph summary, intended to be read by other experts of the same field. In all four categories, the experts aim at questions for which a limited number of articles (min. 10, max. 60) are retrieved through PubMed queries. Questions which are controversial or that have no clear answers in the literature are avoided. Moreover, all questions are related to the biomedical domain. For example, in the case of the following two questions:

Q₁: Which are the differences between Hidden Markov Models (HMMs) and Artificial Neural Networks (ANNs)?

Q₂: Which are the uses of Hidden Markov Models (HMMs) in gene prediction?

Although HMMs and ANNs are used in the biomedical domain, *Q₁* is not suitable for the needs of BioASQ, since there is not a direct indication that it is related to the biomedical domain. On the other hand, *Q₂* links to “gene prediction” and is appropriate.

Step 2: Relevant concepts. A set of terms that are relevant to each question is selected. The set of relevant terms may include terms that are already mentioned in the question, but it may also include synonyms of the question terms, closely related broader and narrower terms etc. For the question “*Do CpG islands colocalise with transcription start sites?*”, the set of relevant terms would most probably include the question terms “*CpG Island*” and “*transcription start site*”, but possibly also other terms, like the synonym “*Transcription Initiation Site*”.

Step 3: Information retrieval. Using the selected terms, the BioASQ annotation tool allows the experts to issue queries and retrieve relevant articles through PubMed. More than one query may be associated with each question and each query can be enriched with the advanced search tags of PubMed. The search window (Figure 5) allows selecting information that is necessary to answer the question. One of the main powers of the annotation tool is that it implements interfaces to different data sources of different types, i.e., unstructured, semi-structured or structured. Given that we cannot expect domain experts to be familiar with Semantic Web standards, such as RDF, the annotation tool also implements an innovative natural language generation method that converts RDF into natural language. The iterative improvement of the annotation tool has led to a framework that is widely accepted by the BioASQ biomedical expert team. Interestingly, a study of the queries used by different experts to answer the same questions made clear that indeed “many roads lead to Rome”, i.e. different experts will use different queries for the same question.

“Putative Zinc Finger Protein Binding Sites Are Over-Represented in the Boundaries of Methylation-Resistant CpG Islands in the Human Genome”

“CpG Islands: Starting Blocks for Replication and Transcription”

Table 1. Examples of retrieved articles (only titles shown here, but the annotation tool provides also the abstracts).

Selected results		
C	"acetonitrile"	-
D	Nano-intercalated rhodanese in cyanide antagonism	-
D	Release of sunflower seed dormancy by cyanide: cross-talk with ethylene signalling pathway	-
C	Manihot	-
D	Gene-based microsatellites for cassava (<i>Manihot esculenta</i> Crantz): prevalence, polymorphisms, and cross-taxa utility	-
D	Effect of molasses on nutritional quality of cassava and gliciridia tops silage	-
D	Biofortification of essential nutritional	-

Release of sunflower seed dormancy by cyanide: cross-talk with ethylene signalling pathway

Introduction Cyanide is a compound known to stimulate germination and to release dormancy of seeds of many species (Taylorson and Hendricks, 1973; Roberts and Smith, 1977; Bogatek and Lewak, 1988; Côme et al., 1988; Bethke et al., 2006). Seed dormancy is defined as the property of a seed that prevents its germination in apparently favourable conditions (Finch-Savage and Leubner-Metzger, 2006). Despite the well-described effects of cyanide on germination and dormancy, the cellular bases of its mechanism are poorly understood, and seem moreover to vary from one species to another. Different hypotheses have been proposed to explain the stimulatory effect of cyanide on germination and dormancy (Côme and Corbineau, 1989). According to Taylorson and Hendricks (1973), the cyanohydrin gas could react with L-cysteine to give the β -L-cyanoalanine necessary for the synthesis of arginine and aspartic acids which could be limiting factors for germination. Hagesawa et al. (1994) suggested that this increase in the amino acid pool might also promote germination by decreasing the water potential in embryonic axis. However, other respiratory inhibitors which are not metabolized, such as NaN₃ or Na₂S, have the same effect as KCN in various species (Roberts and Smith, 1977; Côme and Corbineau, 1989). Some studies proposed that the beneficial effect of cyanide on germination might involve the cyanide-insensitive pathway (Esashi et al., 1979, 1981b; Upadhyaya et al., 1983), the pentose phosphate pathway (Roberts and Smith, 1977; Côme and Corbineau, 1989), the glycolysis (Bogatek, 1995) or the hydrolysis of oligosaccharides and their catabolism (Bogatek and Lewak, 1991; Bogatek et al., 1999). Cyanide is also known to interact with reactive oxygen species (ROS) metabolism; it is an inhibitor of Cu/Zn superoxide dismutase (SOD) (Bowler et al., 1992) and catalase (CAT) (Tejera García et al., 2007) and it has been demonstrated to induce oxidative stress and lipid peroxidation in animals (Johnson et al., 1987; Gunasekar et al., 1998). Oracz et al. (2007) recently demonstrated that cyanide could trigger protein oxidation during sunflower seed dormancy alleviation. At last, cyanide might also interplay with the ethylene signalling pathway. Indeed, hydrogen cyanide is a co-product of ACC oxidase, which converts ACC to ethylene (Peiser et al., 1984), and it has been proposed to stimulate ethylene biosynthesis via a feedback effect (Pirrung and Brauman, 1987). Thus Smith and Arteca (2000) demonstrated that the ACC synthase gene ACS6 was activated by cyanide in *Arabidopsis*. However, it is actually not known whether ethylene and cyanide share some molecular components of their downstream transduction pathways. The putative relationship between cyanide and ethylene signalling pathways might be particularly relevant for sunflower seeds, whose dormancy is broken by ethylene (Corbineau et al., 1990). The inability of freshly harvested sunflower seeds to germinate at temperatures below c. 15 °C results from an embryo dormancy which is gradually eliminated during dry storage (Corbineau et al., 1990; Corbineau and Côme, 2003). Embryo dormancy is characterized by the inhibition of radicle extension thus preventing excised embryos to grow (Finch-Savage and Leubner-Metzger, 2006). Little attention has been paid to the

Fig. 6 Screenshot of the annotation tool’s snippet annotation process.

Returning to the example question “Do CpG islands colocalize with transcription start sites?” a query may be “CpG Island” AND “transcription start site”. Some of the articles retrieved by this query are shown in Table 1.

Step 4: Selection of articles. Based on the results of Step 3, the experts select a set of articles that are sufficient for answering the question. Using the annotation tool, they choose among the retrieved list of articles, the ones that contain relevant information to form an answer.

Step 5: Text snippet extraction. Using the articles selected in step 4, the experts mark every text snippet (piece of text) out of the articles selected in Step 4. Snippets can be easily extracted using the annotation tool (Figure 6) and may answer the question either fully or partially. A text snippet should contain one or more entire and consecutive sentences. If there are multiple snippets that provide the same (or almost the same) information (in the same or in different articles), all of them are selected. Examples of relevant snippets are shown in Table 2.

Step 6: Query revision. If the expert judges that the articles and snippets gathered during steps 2 to 5 are insufficient for answering the question, the process can be repeated. The articles that the expert has already selected can be saved before performing a new search, along with the snippets the expert has already extracted. The query can be revised several times, until the expert feels that the gathered information is sufficient to answer the question. At the end, if the expert judges that the question can still not be answered adequately, the question is discarded.

“A common explanation for the G + C rise that is seen here in the mammalian profile in the proximity of the TSS is the presence of CpG islands.”

“Above we have made the remark that the G + C rise in mammals and maybe generally in vertebrates is probably caused by the higher number of CpG dinucleotides in the promoter region.”

Table 2. Examples of relevant snippets.

“Yes. It is generally known that the presence of a CpG island around the TSS is related to the expression pattern of the gene. CGIs (CpG islands) often extend into downstream transcript regions. This provides an explanation for the observation that the exon at the 5' end of the transcript, flanked with the transcription start site, shows a remarkably higher CpG density than the downstream exons.”

Table 3. Example of ideal answer.

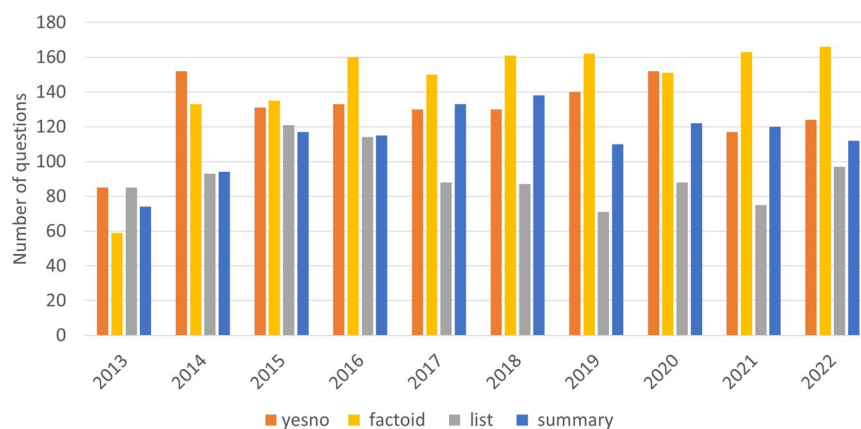


Fig. 7 Distribution of types of questions per year.

Step 7: Exact answer. In steps 2 to 6, the expert identifies relevant material for answering the question. Given this material, the next step is to formulate the actual answer. For a yes/no question, the exact answer is simply “yes” or “no”. For a factoid question, the exact answer is the name of the entity (e.g., gene, disease) sought by the question; if the entity has several synonyms, the expert provides, to the extent possible, all of its synonyms. For a list question, the exact answer is a list containing the entities sought by the question; if a member of the list has several synonyms, the expert provides again as many of the synonyms as possible. For a summary question, the exact answer is left blank. The exact answers of yes/no, factoid, and list questions should be based on the information of the text snippets that the expert has selected, rather than personal experience.

Step 8: Ideal answer. At this final step, the expert formulates what we call an *ideal answer* for the question. The ideal answer should be a one-paragraph text that answers the question in a manner that the expert finds satisfactory. The ideal answer should be written in English, and it should be intended to be read by other experts of the same field. For the example question “Do CpG islands colocalise with transcription start sites?”, an ideal answer might be the one shown in Table 3. Again, the ideal answer should be based on the information of the text snippets that the expert has selected, rather than personal experience. The experts, however, are allowed (and should) rephrase or shorten the snippets, order or combine them etc., in order to make the ideal answer more concise and easier to read.

Notice that in the example above, the ideal answer provides additional information supporting the exact answer. If the expert feels that the exact answer of a yes/no, factoid, or list question is sufficient and no additional information needs to be reported, the ideal answer can be the same as the exact answer. For summary questions, an ideal answer must always be provided.

Figure 7 presents the distribution of questions created each year of the challenge. Over the years, there is an increase in the number of factoid questions, and a decrease in the number of list questions. The possible reason is that it is more difficult to find the material (i.e. articles and snippets) that are sufficient for answering a factoid question than a list question. Table 4 presents the different versions of the BioASQ-QA dataset, including the number of questions, and the average number of documents and snippets. Each version of the training dataset enriches its previous version with the new questions created the respective year.

During the years that BioASQ has been running, three significant changes have taken place, in response to feedback obtained by the experts and the challenge participants:

Versions of data	Size (cumulative)	Documents (average)	Snippets (average)
2013	310	14.28	18.71
2014	810	13.45	13.30
2015	1,307	13.00	17.86
2016	1,799	11.86	20.38
2017	2,251	12.01	14.76
2018	2,747	11.14	13.91
2019	3,243	10.15	12.92
2020	3,742	9.43	12.33
2021	4,234	9.22	12.24
2022	4,721	8.58	11.36

Table 4. The different versions of the BioASQ-QA dataset, as it has evolved over the years of the challenge. BioASQ 1 produced only 10 questions, as the first round of the challenge acted as a proof-of-concept. These 10 questions have been incorporated into the BioASQ 2 set (2013).

Ideal answers:

Sp1 binds to a GC-rich sequence element containing the decanucleotide consensus sequence 5'-(G/T)GGGCGG(G/A)(G/A)(C/T)-3' (GC box element) in double stranded DNA (dsDNA). Gel shift competition studies and DNase I footprinting analyses revealed that Sp1 specifically interacts with the CACCC motif.

We have previously shown that mutations in the GGAA core motif of the Ets1 binding site, EBSI, or deletion of EBSI, reduced basal and Tax1 transactivation of the PTHrP P2 promoter. Adipocyte amino acid transporter is induced during the 3T3-L1 preadipocyte differentiation process. Site-specific mutations in the CACCC motif decreased promoter

Information recall:

Information precision:

Information repetition:

Readability:

1 2 3 4 5

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Information repetition:

Readability:

1 2 3 4 5

Fig. 8 Assessment tool for evaluating system answers. The gold standard answer is at the top.

Filed	Type	Content
id	String	A unique identifier of the question. E.g. "52bf1b0a03868f1b06000009"
body	String	The question body in English. E.g. "What is the mode of inheritance of Wilson's disease (WD)?"
type	String	The question type in English. One of "yesno", "factoid", "list" or "summary"
documents	Array of Strings	List of relevant article URLs. E.g. ["https://www.ncbi.nlm.nih.gov/pubmed/838566",...]
snippets	Array of JSON Objects	List of relevant snippets. E.g. [{"offsetInBeginSection":122,"offsetInEndSection":272,"text":"The disease...","beginSection":"abstract","document":"http:...","endSection":"abstract"},...]
concepts	Array of Strings	List of relevant concept URLs. E.g. ["https://www.disease-ontology.org/api/metadata/DOID:893",...]
triples	Array of JSON Objects	List of relevant triples. E.g. [{"p":"http.../name", "s":"http.../diseases/1198", "o":"Wilson_disease"},...]
ideal_answer	Array of Strings	List of ideal answers to the question in English. E.g. ["WD is an autosomal recessive disorder",...]
exact_answer not available in summary questions	Depends on the type of the question	For <i>yesno</i> : A String ("yes" or "no") For <i>factoid</i> : An array of Strings, synonyms of the answer. E.g. ["CaM kinase II", "CAMK2"] For <i>list</i> : An array of arrays of Strings with synonyms of each element of the answer. E.g. [{"Triadin","TrD"}, ["Calsequestrin", "CASQ",...],...]

Table 5. JSON format of the BioASQ-QA benchmark dataset.

- Since BioASQ 3 (2015), the focus of the experts is only on relevant articles and their contents. In other words, the experts do not provide relevant concepts or statements, as it was found cumbersome and led to questionable results. Nevertheless, concepts are included in the gold dataset, as they are added by the systems and assessed by the experts in the assessment phase).
- Since BioSQ 4 (2016) only a sufficient set of articles, that allow the answer to be found with confidence, is requested by the experts. This is again in contrast to earlier years, where the experts were asked to identify all relevant articles; something that proved to be unrealistic. Again, if the participating systems retrieve more

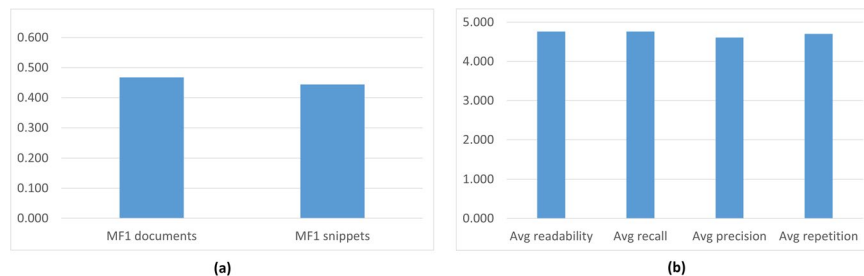


Fig. 9 Inter-annotator agreement: (a) Scores of the snippets and the documents retrieved by the additional expert, compared to the original one; and (b) average manual scores of the expert ideal answers.

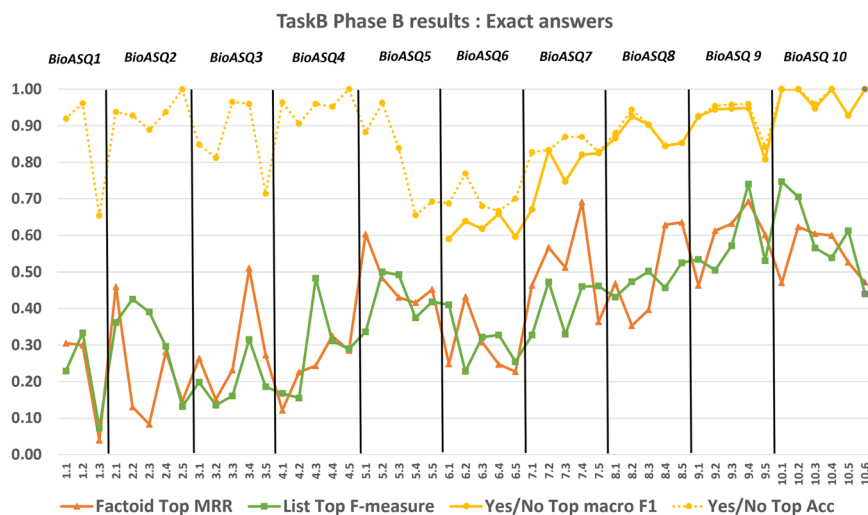


Fig. 10 The performance achieved by systems in exact answer generation, across different years of the BioASQ challenge. For each test set the performance of the best performing system (Top) is presented based on the official evaluation measures. Since BioASQ6 the macro-averaged F1 score (macro F1) is the official measure for Yes/No questions, but accuracy (Acc), the former official measure, is also presented.

relevant documents, not identified in the annotation phase by the experts, these are added in the gold dataset, during the assessment phase.

- In early versions of the challenge, we considered using full-text articles from PubMed Central (PMC). Given the small percentage of the overall literature that appears in PMC, since BioASQ 4 (2016) we decided to restrict the challenge to article abstracts only.

Assessment. Following each round of the challenge, the answers of the participating systems are collected and assessed. Exact answers can be assessed automatically against the golden answers provided by the experts during the annotation phase. However, the ‘ideal’ answers are assessed manually by the experts. In fact each expert gets to assess the answers to the questions they have created, in terms of *information recall* (does the ‘ideal’ answer reports all the necessary information?), *information precision* (does the answer contain only relevant information?), *information repetition* (does the ‘ideal’ answer avoid repeating the same information multiple times? e.g., when sentences of the ‘ideal’ answer that have been extracted from different articles convey the same information), and *readability* (is the ‘ideal’ answer easily readable and fluent?). A 1 to 5 scale is used in all four criteria (1 for ‘very poor’, 5 for ‘excellent’).

The assessment tool is designed to be a companion to the annotation tool and is implemented by reusing most of its functionality. The tool can also be used to perform an inter-annotator agreement study. In that case, domain experts are provided with answers generated by other (anonymous) domain experts and are asked to evaluate them.

The design of the interface is such that the users can always see the answers/annotations only to questions that they are asked to review (Figure 8). Moreover, the interface can adapt to different question types, by showing different answering fields for each of them. Finally, all information sources that were used to answer the question can also be reviewed. By these means, domain experts can perform an informed assessment.

The assessment tool plays a key role in the creation of the benchmark and the quality assurance of the results generated by the experts during the BioASQ challenges. Moreover, the assessment tool allows the experts to improve their own gold answers and associated material, based on the answers provided by the systems. In

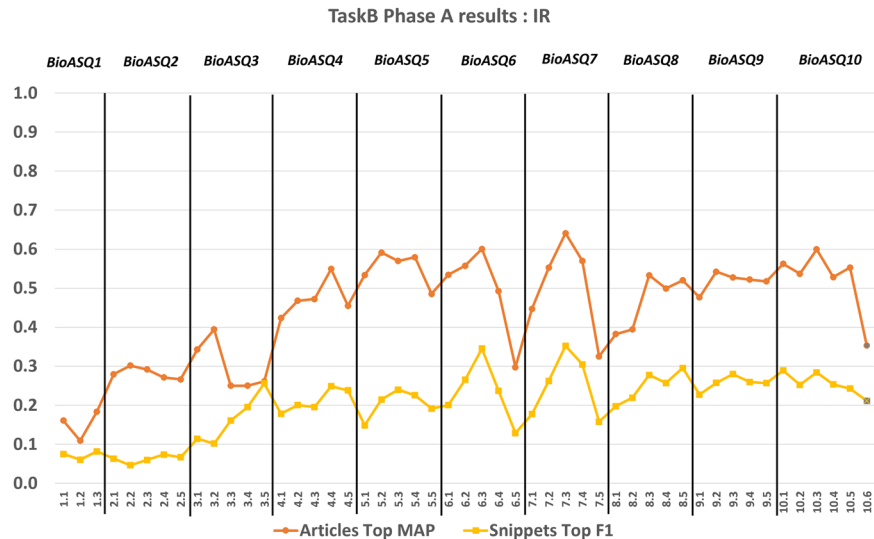


Fig. 11 The performance achieved by systems in the information retrieval part of Task B, across different years of the BioASQ challenge. For each test set the performance of the best performing system (Top) is presented, based on the official evaluation measures.

particular, the experts revise the documents and snippets returned by the systems, and enrich the gold answers with material identified by the systems. This leads to an improvement of the benchmark datasets that are provided publicly.

Data Records

The dataset is available at Zenodo¹⁵ and follows the *JSON* format. Specifically, it contains an array of questions, where each question (represented as an object in the *JSON* format) is constructed as shown in Table 5.

Technical Validation

Improving the state-of-the-art performance. The participation of strong research teams in the BioASQ challenge has helped to measure objectively the state-of-the-art performance in biomedical question answering^{16–18}. During BioASQ this performance has improved (Figures 10, 11, 12). It is particularly encouraging that the BioASQ biomedical expert team have assessed the ideal answers provided by the participating systems as being of very good quality. The average manual scores are above 80% (above 4 out of 5 in Fig. 12). Still, there is much room for improvement and the future challenges of BioASQ, as well as the benchmark datasets that it provides, will hopefully push further towards that direction.

Based on the evaluation of the participating systems from the experts, one very interesting result is that humans seem satisfied by “imperfect” system responses. In other words, they are satisfied if the systems provide the information and answer needed, even if it is not perfectly formed.

Inter-annotation agreement. An inter-annotation agreement evaluation has been conducted in order to evaluate the consistency of the dataset. To this end, during the first six years, a small subset of the created questions, was given to other experts, in order to compare the different formulated answers. The pairs of experts answer the exact same questions. As each of them uses their own queries, they get a different list of possible relevant documents. The latter leads to the selection of different documents and snippets to answer the questions, which leads to low mean F1 score (<50%) (Figure 9a). Nevertheless, in the formulated ideal answers, there is a high agreement between the experts (Figure 9b). In other words, they reach the same or very similar answers, but following different paths.

Another important point is that the BioASQ challenge, as well as the environment in which it takes place, evolve. One consequence of this is the changes that we had to make in the data generation process in response to feedback from the experts and the participants. Additionally, the evolution of vocabularies and databases cause complications. For example, each year’s data are annotated with the current version of the MeSH hierarchy, which is updated annually. In addition, only the articles of the current year of annotation are used for formulating the answers, while articles that will appear in the future may also be of relevance. These are issues that we need to handle and adapt to them, in order to have real-life, useful challenge and relevant dataset.

The BioASQ challenge will continue to run in the coming years, and the dataset will be further enriched with new interesting questions and answers.

Usage Notes

Up to date guidelines and usage examples pertaining the dataset can be found in: <http://participants-area.bioasq.org/>

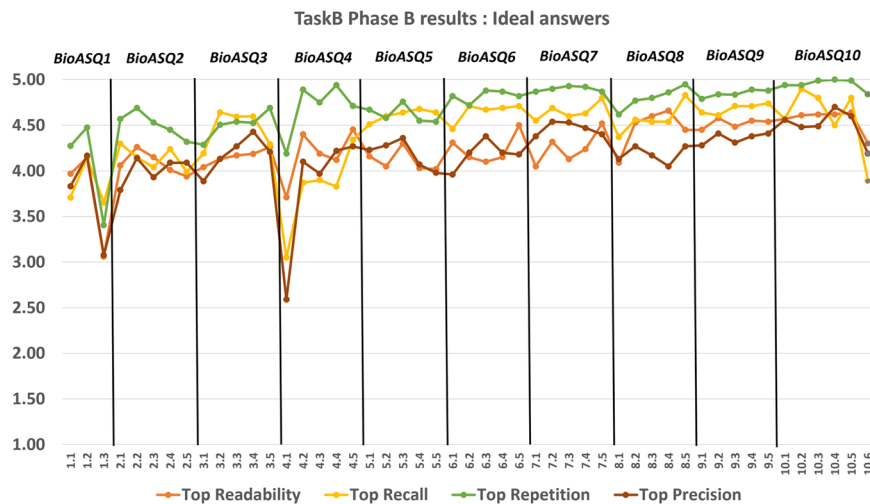


Fig. 12 The performance achieved by systems in the ideal answer generation part of Task B, across different years of the BioASQ challenge. For each test set the performance of the best performing system (Top) is presented based on the official evaluation measures. In BioASQ 4, the performance was lower due to the fact that it was the first time BioASQ run after the end of the EU-funded project, and the top teams in ideal answer generation started participating after the second batch of the same year.

Code availability

BioASQ has created a lively ecosystem, supported by tools and systems that facilitate the creation of the benchmarks. All software is provided with open-source licenses (<https://github.com/BioASQ>). In addition, the data produced are open to the public¹⁵.

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Author contributions

G.P. and A.K. originated the BioASQ challenge and dataset creation. All authors participated in the collection of the data, the development of the annotation and the evaluation tools, and validated the data. A.K. and A.N. drafted the manuscript. All authors reviewed the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

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