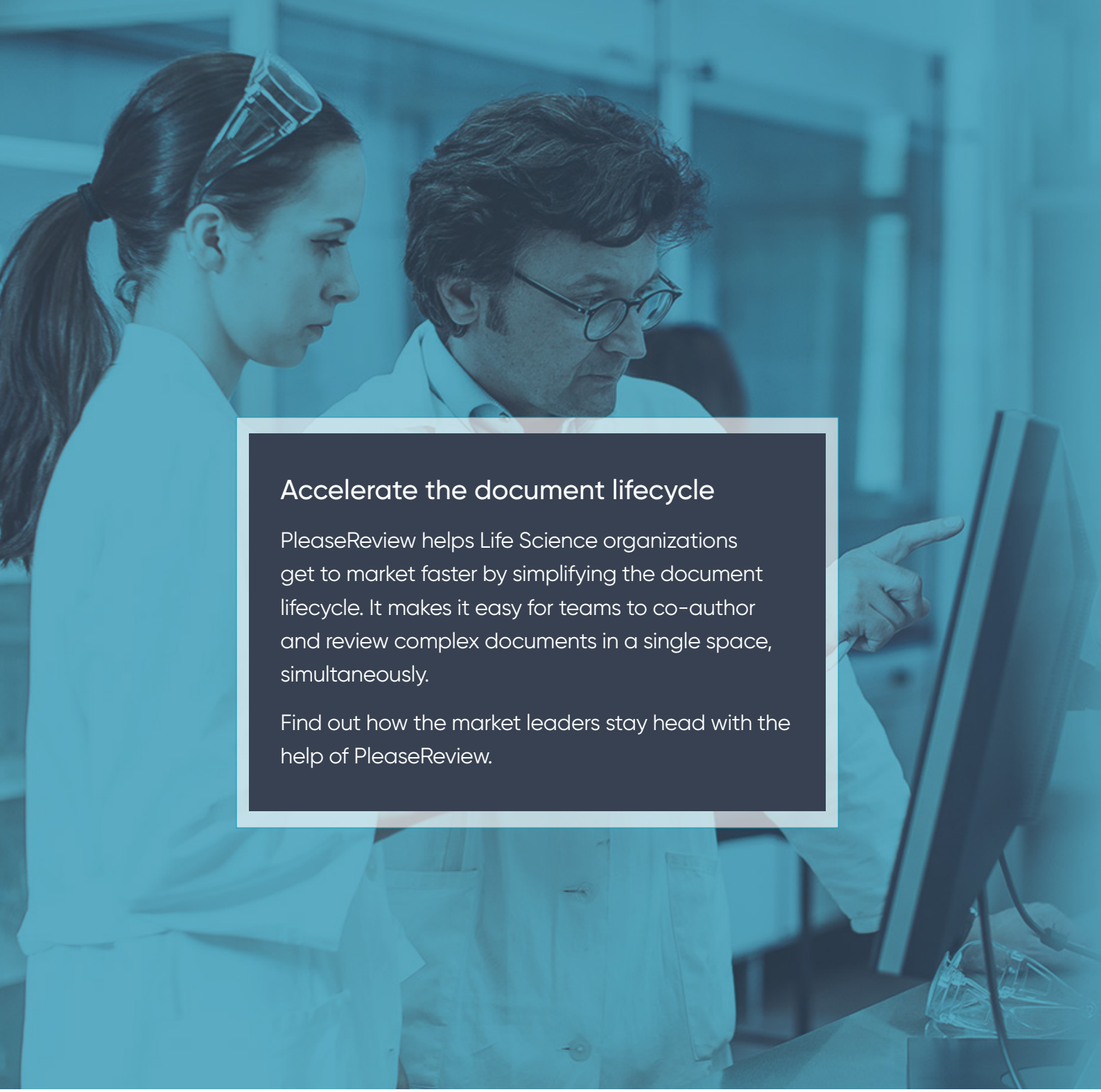
Ideagen

PleaseReview Success Stories

Discover the Business Case for PleaseReview



Accelerate the document lifecycle

PleaseReview helps Life Science organizations get to market faster by simplifying the document lifecycle. It makes it easy for teams to co-author and review complex documents in a single space, simultaneously.

Find out how the market leaders stay head with the help of PleaseReview.

Trusted by many well-known Life Science brands

Lexicon pharmaceuticals

VERTEX

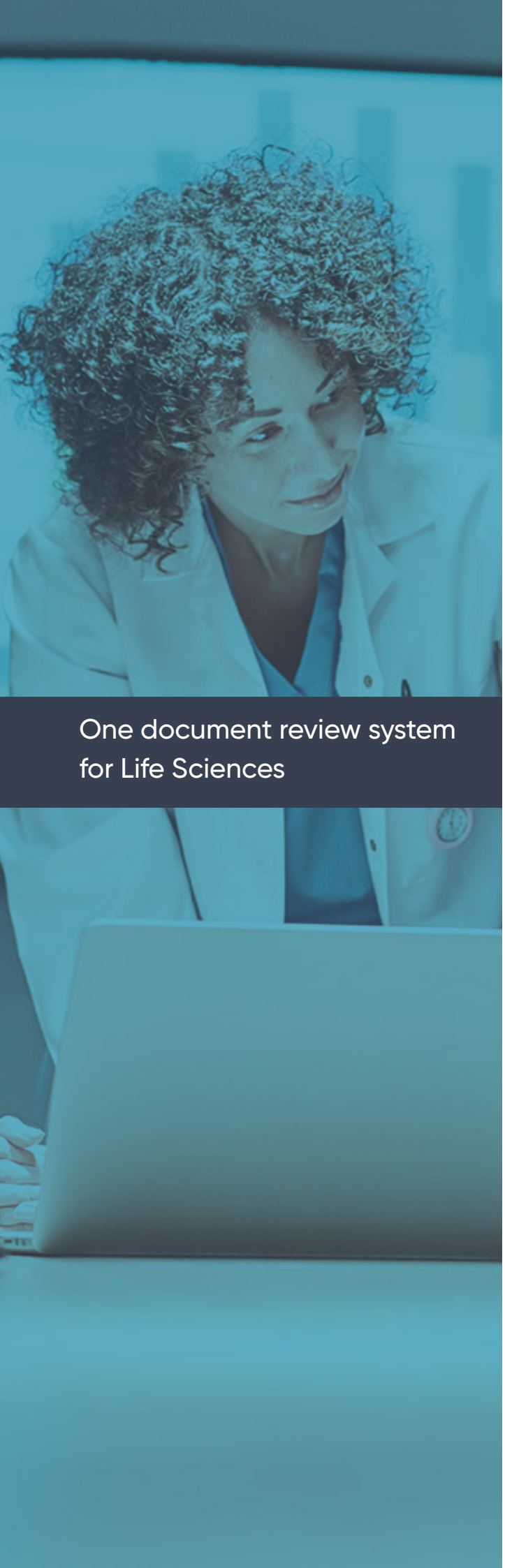
ucb Pharma

Alnylam® PHARMACEUTICALS

ACORDA® THERAPEUTICS

pharmacyclics® An AbbVie Company

GALDERMA



One document review system for Life Sciences

Support all your teams that collaborate on highly sensitive regulatory documents:

- Investigational new drug
- New drug applications
- FDA clinical submissions
- Emergency use authorization
- Investigator's brochures
- Medical device reports
- Clinical research
- Medical writing

In response to the COVID-19 crisis, the top 5 approved vaccines were developed in record time with the help of PleaseReview.

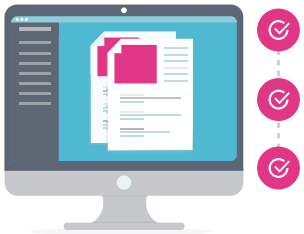
PleaseReview brings commercial value to your business and efficiency to your teams.

Developing a new drug or medical device can be a lengthy endeavor and cost billions of dollars. When it takes an average of 12 years to progress a drug from discovery to market, you do not want slow and outdated processes to be the reason your competitors beat you to the finish line. That's why so many life science companies use PleaseReview to speed up the document collaboration and review process.

TOP BENEFITS

Collaborate on the same documents simultaneously

Keep sensitive data secure



Comply with regulatory and industry standards

> We have saved literally hundreds of hours over a year.
>
> **Kristi Boehm, Head of Medical Writing – Lexicon Pharmaceuticals**

TIME SAVINGS

Using existing review tools

Using PleaseReview

This demonstrates an elapsed time saving of 9 days for a single document

Reviewer savings and Author 'work time' savings by halving review cycles required

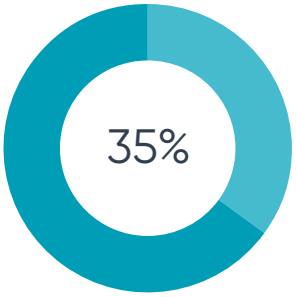
For each document reviewed there is an overall **6 person hours saving**, representing nearly a person day for every document.

> PleaseReview has been a great success in our company and our users have been happy with how easy the software is to use. I continue to receive positive feedback from them.
>
> **Regulatory Affairs Manager**

> We will be advancing our R&D and clinical programs moving forward and PleaseReview will be the tool we use throughout this.
>
> **Thomas Class, Head of Regulatory Affairs – Translate**

COST SAVINGS

PleaseReview saved a large scale project at a major company over **$150 per hour = $5000 per document**



On average, PleaseReview is proven to provide savings of **35%** compared with manual processes

Proven to manage even the heaviest document workload:

150+
documents

3000+
comments

50+
reviewers

SUCCESS STORY

Translate Bio

The Challenge

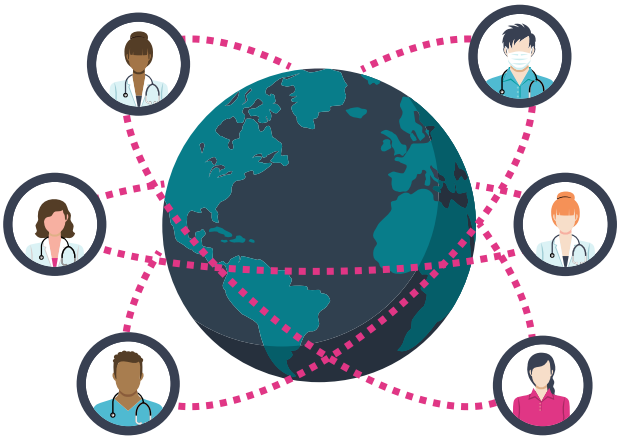
- Sizeable teams reviewing large documents
- Comments and updates lost or overlooked
- Sharing documents by email causing version confusion

> Before using PleaseReview, the Contract Medical Writers were responsible for sending out documents for review via email and then collating comments back into one document, which was an archaic way of doing things.

The Solution

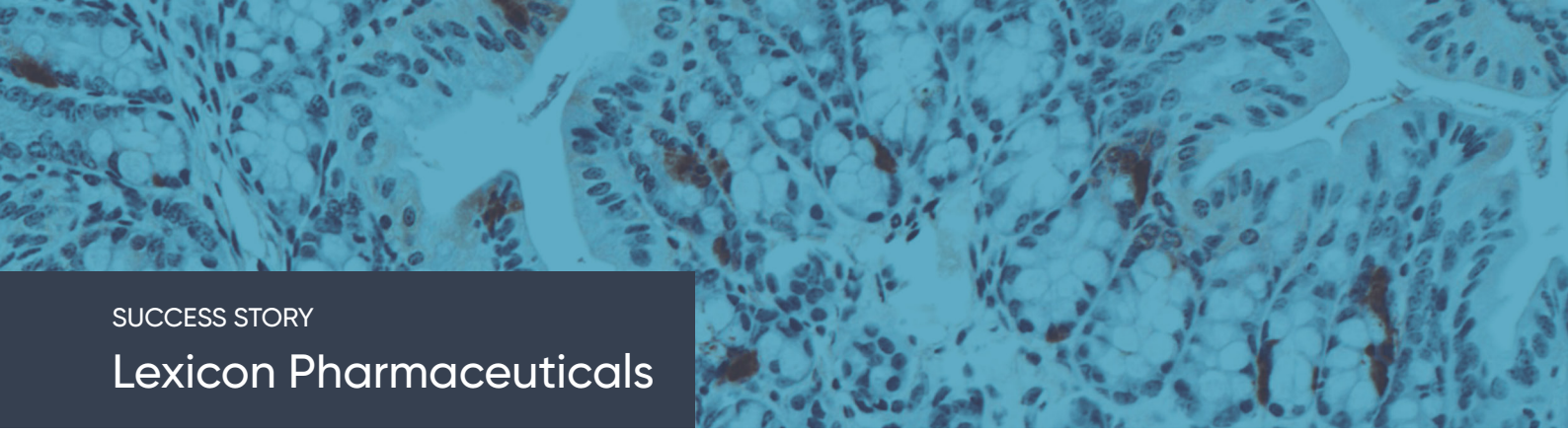
Translate Bio were able to include large teams on the review process of lengthy documents, which was key to their ability to efficiently review and approve quickly. By speeding up their entire review cycle, document owners were able to easily consolidate comments. Where they used to have multiple documents and differing versions, PleaseReview provided a single version with teams collaborating all in one place.

Using PleaseReview has allowed Translate Bio to seamlessly collaborate with regulatory authorities around the world.



See the full story.

> PleaseReview provides us with a stable review tool that is reliable and straightforward. I think, particularly for pharmaceutical companies, it is a valuable tool. It really helps us to get through the process of a document collaboration review in a much more orderly manner.
>
> **Thomas Class, Head of Regulatory Affairs – Translate Bio**



SUCCESS STORY

Lexicon Pharmaceuticals

The Challenge

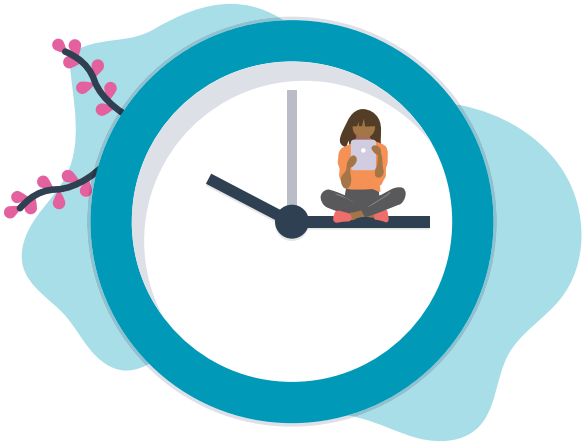
- Up to 60 people in a review
- Multiple drug trials in progress
- Lack of oversight and control

> Before PleaseReview, everyone reviewed in a vacuum, without being able to see anyone else's edits or comments. Conflicting edits and dispute resolution were a nightmare.

The Solution

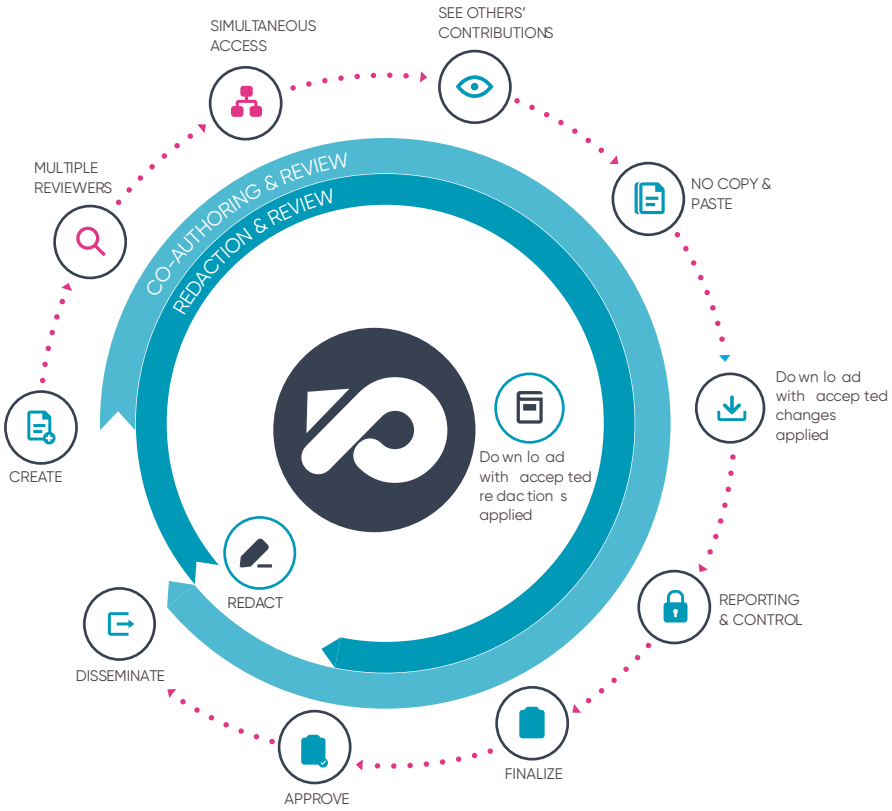
The medical writing team at Lexicon Pharmaceuticals has dramatically increased its productivity with real-time document collaboration. Projects can be turned around much faster because of the time saved in collating, editing, and incorporating changes into document reviews.

Lexicon Pharmaceuticals saved hundreds of hours by reducing document review times with PleaseReview.



See the full story.

> I am 100% delighted with PleaseReview. It has freed me from days of comment incorporation and playing referee!"
>
> **Kristi Boehm, Manager for Medical Writing, Lexicon Pharmaceuticals**



Accelerating to market

PleaseReview is used widely by Medical Writing, Regulatory, Clinical, Quality, and other professionals within the Life Sciences sector.

As documents become more complex, more people need to be involved in their creation and review. The more people that are involved, the harder it is to understand what changes have been suggested and by whom.

Collaborators and reviewers can seamlessly view, comment, edit, and approve reports, submissions, and publications all in one place without the worry of missing any contributions.

PleaseReview helps you to control and manage all aspects of document collaboration. The web-based software solution helps you:

- Work more effectively with teams and stakeholders.
- Have better control of the document – keeping all updates and comments within a single version.
- Cut the risk of document error.
- Keep a full audit by automatically creating a complete report of all activity on the review.
- Secure sensitive data throughout the document lifecycle.

> We have been able to almost eliminate the amount of time authors spend collating comments into the next version and are now far more efficient about resolution of conflicting reviewer comments.
>
> **Regulatory Affairs Manager**

About Ideagen

Ideagen plc develops software that helps to solve complex problems in highly regulated industries including life sciences, healthcare, financial services, government, manufacturing, and utilities.

With more than 20 years of experience and success in the global regulatory environment, we have earned our customers' trust by building long-term relationships to meet and exceed their risk, compliance, and audit management goals.

PleaseReview helps to cut product development time and costs with a simple way to manage collaboration on regulatory documents.

Join the industry standard and accelerate your journey to market.

Book a demo

Ideagen

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