

FACET LIFE SCIENCES

PRESENTS

GEMSTONE

The Facet Life Sciences team has decades of experience working with small development teams. We understand goals of small teams and have developed Gemstone, our electronic document management solution, as the simple, secure and compliant document management tool of the trade.

Document management for small biopharmaceutical teams can open the company to unintended risks. Choosing a document management solution should streamline processes while protecting valuable content. If it doesn't, it isn't the right tool. Facet Life Sciences is redefining document management for small teams. Our practice-based approach is driven by experience, expertise and a steadfast commitment to supporting small, innovative teams with right-sized technology solutions.



FACET

GEMSTONE



OUR PACKAGES

SIMPLE
CUSTOMIZEABLE

The rationale behind Facet Gemstone is to offer a simple electronic document management solution for small teams that supports the complex development document lifecycle. Facet Gemstone is the essential productivity tool for life science teams. It enables content driven processes with simple, secure and compliant features.

DISCOVERY & EARLY DEVELOPMENT



Secure your asset

You are a scientist and/or CEO. You have a valuable new molecule, device, or delivery modality: a rare gem. You have proprietary background materials and early development information that require secure storage and controlled sharing. You need a secure, reliable and compliant tool to protect your asset. Does this sound like you?

IND & CLINICAL DEVELOPMENT



Collaboration, clinical testing and growth

You are ready to get your rare gem into and through clinical development. You need to share documents with a lead scientist or a contract research group. You keep control of your documents and data while enabling collaboration with key team members. Does this sound like you?

MARKETING APPLICATION SUBMISSIONS



Prepare and organize documentation

Your gem is being polished to perfection. You are preparing to submit a marketing application. You can finalize documents while simultaneously organizing for health authority submission and preparing for your submission publishing partner in a safe and secure environment. Does this sound like you?

DUE DILIGENCE REVIEW FACILITATION



Facilitate due diligence reviews

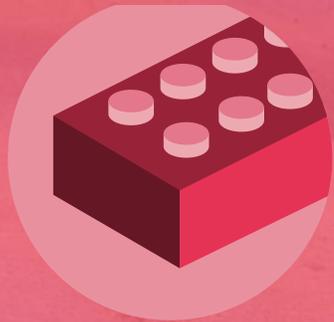
At any stage in development, you may want to share information with investors or position your company and/or asset for acquisition. You've also got full control to grant read only access for the due diligence or review process as needed. Does this sound like you?

DOCUMENT MANAGEMENT IS
NEEDED AT EVERY STAGE OF DRUG
DEVELOPMENT



NEED
SOMETHING
DIFFERENT?

We understand that every business is different, and customization is available.



SIMPLE

In order for a solution to be considered a “tool of the trade”, it must be easy to understand, easy to use on a regular basis and require minimal training, customization, and ongoing IT support. Gemstone provides a simple user interface that requires a one-time, less than one-hour training. The intuitive nature of the interface speeds up time to productive daily use. Simple upload and versioning functionality streamline document review and approval processes.



SECURE

Gemstone addresses security at multiple levels within the product. The product provides a secure connection via the web, using two-step authentication from the desktop to the cloud. Our technology partners have achieved multiple security certifications including ISO 27001, SSAE 16, have earned a TRUSTe Certified Privacy Seal and are a part of the Cloud Security Alliance.



COMPLIANT

Gemstone is 21 CFR, Part 11 compliant. A human readable audit trail is produced for all electronic records managed within the system. The audit trail identifies who changed the record, what they changed and when they changed it. Part 11 compliance is essential for R&D documentation that will be included in a regulatory submission. Prospective partners and investors will appreciate the assurances that the documentation has been created in a controlled environment.

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