Legility

Prescription for Success Experienced. Efficient. Secure.

The pain-free way to manage pharmaceutical document review projects starts with Legility

Since 1991, Legility has managed the eDiscovery and document review process for global pharmaceutical brands and medical device companies. Our team of experienced patent attorneys draws on highly specialized knowledge, including USPTO patent agent certifications and engineering and bio-science degrees, to support the most sophisticated legal matters.

Experience Counts

There are document reviews. And there are *pharmaceutical* document reviews.

We understand the distinction.

That's why the foundation of our Pharmaceutical Patent Managed Review practice group is centered around people — assembling a team of patent bar-certified attorneys and subject matter experts with direct experience handling:

- DOJ/FTC investigations
- Mergers and acquisitions
- Hatch-Waxman litigations
- Product liability and mass tort actions

Our proprietary Hatch-Waxman (HW) training program immerses our attorneys in the processes and specific issues related to HW matters, including patent fundamentals, the NDA approval process and patenting processes, and chemistry and formulation fundamentals.

Pharmaceutical Document Review Benefits

- ✓ Reduced lead and response times
- ✓ Experienced staff, including patent attorneys and scientists
- ✓ Measurable cost reductions
- ✓ Minimize the need for outside counsel, reducing total project costs
- ✓ Centralized data collection/ storage minimizes data security risks

More than 160 of our attorneys have completed the training, which includes a combination of lectures, computer-based training, and personalized study.

Language Support

Recognizing that many pharmaceutical matters involve multi-national touchpoints, we provide direct support for *every* major language — *not translation services*, but a thorough grasp of the nuances of the languages associated with every project.

Efficiencies That Make (Dollars and) Sense

Our pharmaceutical document review process is results-driven, with experienced staff attorneys who are familiar with technical issues and are able to sharply reduce lead times. Their expertise helps reduce data sets quickly and accurately, allowing them to focus on the prioritization of relevant documents.

They employ redaction techniques that accelerate responsiveness, adopting a collaborative approach that continually captures metrics that drive efficiencies and client outcomes.

Safe and Sound

Recognizing the sensitivity of pharmaceutical-related legal matters, we have implemented rigorous security controls that protect and preserve client data.

Our secure, centralized database stores information in a single, secure platform. Rather than disseminate information to each attorney working on a case, performing piecemeal security vetting on each, along with their endpoints, *we bring the counsel to the data*, applying enhanced security protocols that control their level of access.

The result is redactions that are properly implemented and managed, and personally identifiable information (PII) that remains private and secure.