

## **A Team of Nonclinical Safety Experts**

## Asking the Right Questions at the Right Time to Make Safer, More Effective Drugs

With our collective experience of over 400 years in drug discovery and development, we have the expertise to design a high quality, cost-effective nonclinical safety package that perfectly aligns with your project. Early consideration of potential safety liabilities are an important part of every drug discovery project.

### A Skilled Team Providing Support From Start to Finish

- Develops a balanced view of potential safety risks, directing you towards higher quality lead molecules
- Provides study monitoring, CRO selection and study design
- Includes multiple drug modalities and discliplines including genetic toxicology, safety pharmacology, DMPK, reproductive and juvenile toxicity and investigative toxicology

"The support that we received from the ApconiX team has been excellent and they have helped us both in developing our nonclinical safety thinking and in providing direct support for the clinical trial application. We're looking forward to continuing our collaboration as we progress our exciting new medicines into later-stage development and hopefully the clinic"

## Safety risks are the leading cause of attrition in drug discovery and development.

### A well designed and delivered toxicology package can make the difference between success and failure

#### **Drug Discovery**

Early consideration of potential safety liabilities from the drug target of lead molecules should form part of every discovery project. We will work with you to develop a balanced view of potential safety risks, directing you towards higher quality lead molecules and a greater chance of long term success.



#### **Drug Development**

The regulatory GLP studies that underpin the clinical programme should be individually designed to address the scientific questions posed by the molecule, the indication and intended clinical use. Working in partnership with you, we will build a tailored, cost effective nonclinical safety programme that enables the rapid execution of your clinical and regulatory development strategies.

#### **Due Diligence**

Promising compounds are bought, sold and partnered at all stages of the drug development process. Our experts have conducted many Due Diligence reviews and can give you the assurance you need to make a commercially sound decision.



#### **Nonclinical Study Monitoring**

It is essential that toxicology staudies are designed appropriately, conducted to the highest standards, and reported with clarity. An independent Study Monitor helps to select the right CRO, and ensure delivery of a quality end product that meets the needs of the project.

"We would not be here without your excellent input and help.
You and ApconiX have been absolutely fantastic. Great
planning and execution.....strong, straightforward
communication.....pragmatic and a strong sense of team"

## Toxicology is a complex, multidisclipinary science that requires integration of diverse skill sets and creative thinking.



### **Meet Some of the Team**



With many years' experience in discovery and development teams, our project toxicologists have the knowledge and expertise to provide high quality, costeffective preclinical safety packages perfectly aligned with your clinical objectives. Working seamlessly within project teams, our toxicologists help define the key safety questions to allow selection of a de-risked clinical candidate and rapid progression of the development program. We can design the overall safety strategy, provide oversight of safety/toxicology studies, author regulatory documents as well as providing support during regulatory agency interactions.

















# Meet Som





pharmacokinetics, pathology, immuno-toxicology, CRO selection and study monitoring, impurity qualification and OEL/PDE calculation. Our team of recognised experts will provide you with the best advice at the right time to make the right decisions for your project.









