

**TOXIBAN- activated charcoal suspension**  
**LLOYD, Inc. of Iowa**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**ToxiBan®**

**INDICATION:**

For use as a decontamination agent via adsorption of organic chemical toxicants in domestic animals.

In all cases of suspected poisonings, consult your veterinarian for complete diagnosis and specific treatment.

**DOSAGE AND ADMINISTRATION:**

Give orally 10 to 20 mL per kg body weight (5 to 10 mL per lb) by causing the animal to consume the calculated dose. The product may be given as is or mixed with a small amount of cold water. Repeat with the lower dosage every 6 to 8 hours for up to 24 - 48 hours if necessary. Consult a veterinarian should delivery by stomach tube be needed.

**CAUTION:** ToxiBan Suspension should not be given in conjunction with oral medications as the charcoal may adsorb the therapeutic agent. Maintain patient hydration. Monitor for 4 hours after use for signs of hypernatremia (ataxia, tremors, seizures).

**NOTE:** If catharsis is needed, ToxiBan Suspension with Sorbitol should be used initially. **Tests for ethylene glycol must be conducted before ToxiBan Suspension is given to prevent false positive reactions.**

**Adsorptive Power:..... Per mL**

Each mL of ToxiBan Suspension can adsorb at least 55 mg strychnine sulfate.

**SHAKE WELL BEFORE USE**

**KEEP OUT OF REACH OF CHILDREN**

Store at room temperature. Avoid excess humidity.

**Discard within 48 hours after opening.**

**LLOYD, Inc.**

Shenandoah, IA 51601 USA

List No. 1151

Rev. 0623

**PRINCIPAL DISPLAY PANEL - 240 mL Bottle Label**

NDC 11789-105-50

ToxiBan®

Charcoal-Kaolin  
Suspension

**INDICATION:**

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LLOYD®

Net Volume: 240 mL (8.1 fl oz)

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<b>TOXIBAN</b>			
activated charcoal suspension			
<b>Product Information</b>			
<b>Product Type</b>	OTC ANIMAL DRUG	<b>Item Code (Source)</b>	NDC:11789-105
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	<b>ACTIVATED CHARCOAL</b> (UNII: 2P3VWU3H10) (ACTIVATED CHARCOAL - UNII:2P3VWU3H10)	ACTIVATED CHARCOAL	55 [arb'U] in 1 mL
<b>Inactive Ingredients</b>			
	<b>Ingredient Name</b>		<b>Strength</b>

<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>KAOLIN</b> (UNII: 24H4NWX5CO)	

### Product Characteristics

<b>Color</b>	BLACK (BLACK LIQUID)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11789-105-50	240 mL in 1 BOTTLE		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		08/17/2023	

**Labeler** - LLOYD, Inc. of Iowa (962286535)

**Registrant** - LLOYD, Inc. of Iowa (007281942)

### Establishment

Name	Address	ID/FEI	Business Operations
LLOYD, Inc. of Iowa		962286535	PACK, API MANUFACTURE, LABEL, MANUFACTURE

### Establishment

Name	Address	ID/FEI	Business Operations
LLOYD, Inc. of Iowa		007281942	ANALYSIS

Revised: 8/2023

LLOYD, Inc. of Iowa