



# Hy's Law & Drug- Induced Liver Injury (DILI) – Part II

Author:

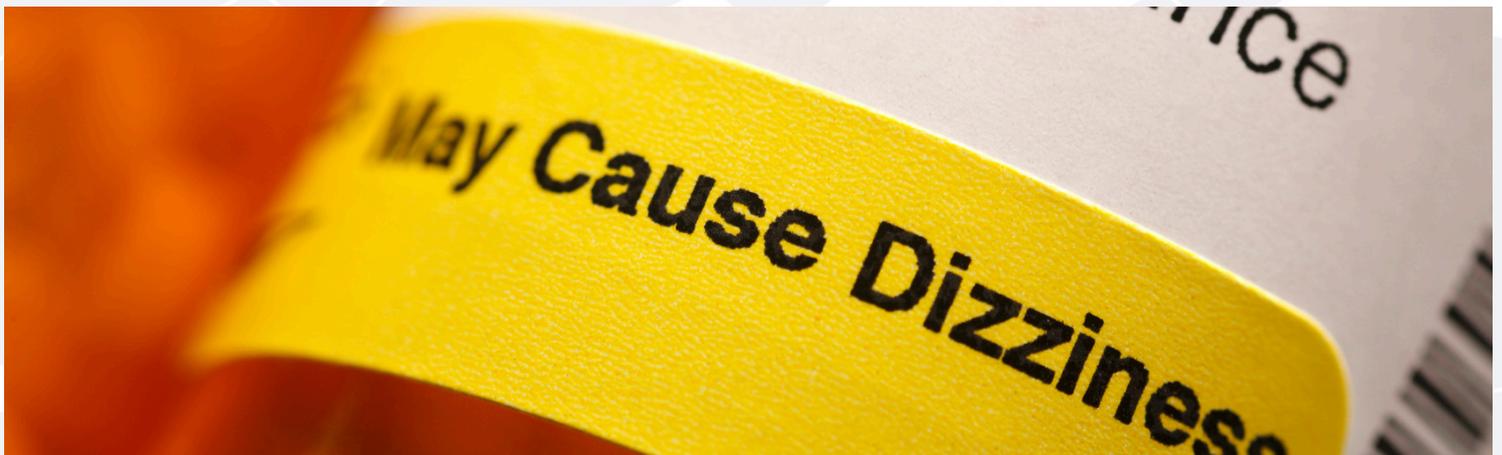
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# Drug-Induced Liver Injury Occurrence

1. Most frequent cause of **acute liver failure** in North America and Europe.
2. No definite causative agent has been attributed in several cases.
3. Underlying mechanisms are still unclear and hence is difficult to predict during drug development.
4. May mimic almost any known type of **liver disease**.
5. Rare yet potentially life-threatening.
6. Key reason for **drugs to fail** to achieve marketing authorization, frequent cause for post-authorization restrictions and product withdrawals.



- Monitoring of standard serum liver tests to detect milder liver injury is the main approach to anticipate a possible DILI risk in Clinical Trials.
- Evaluation of each potential DILI case in clinical trials requires a systematic collection of adequate diagnostic datasets and a rigorous assessment for causality, performed by clinical experts in this area.
- The evaluation of DILI is critical because most drugs that cause severe DILI do so infrequently and usual drug development databases with up to a few thousand subjects exposed to a new drug will not reveal any cases.
- Such databases, on the other hand, may show evidence or signals of a drug's potential for severe DILI, if clinical and laboratory data are properly assessed for evidence of lesser injury, that may not be severe but could predict the ability to cause more severe injuries.

# FDA's eDISH Program for Hepatotoxicity Assessment

**Hy's Law and eDISH Development:** The Hy's Law principle served as the foundation for the FDA's creation of the 'eDISH' software program, designed to **evaluate Drug-Induced Serious Hepatotoxicity**.

## Step-Based Approach:

# 1

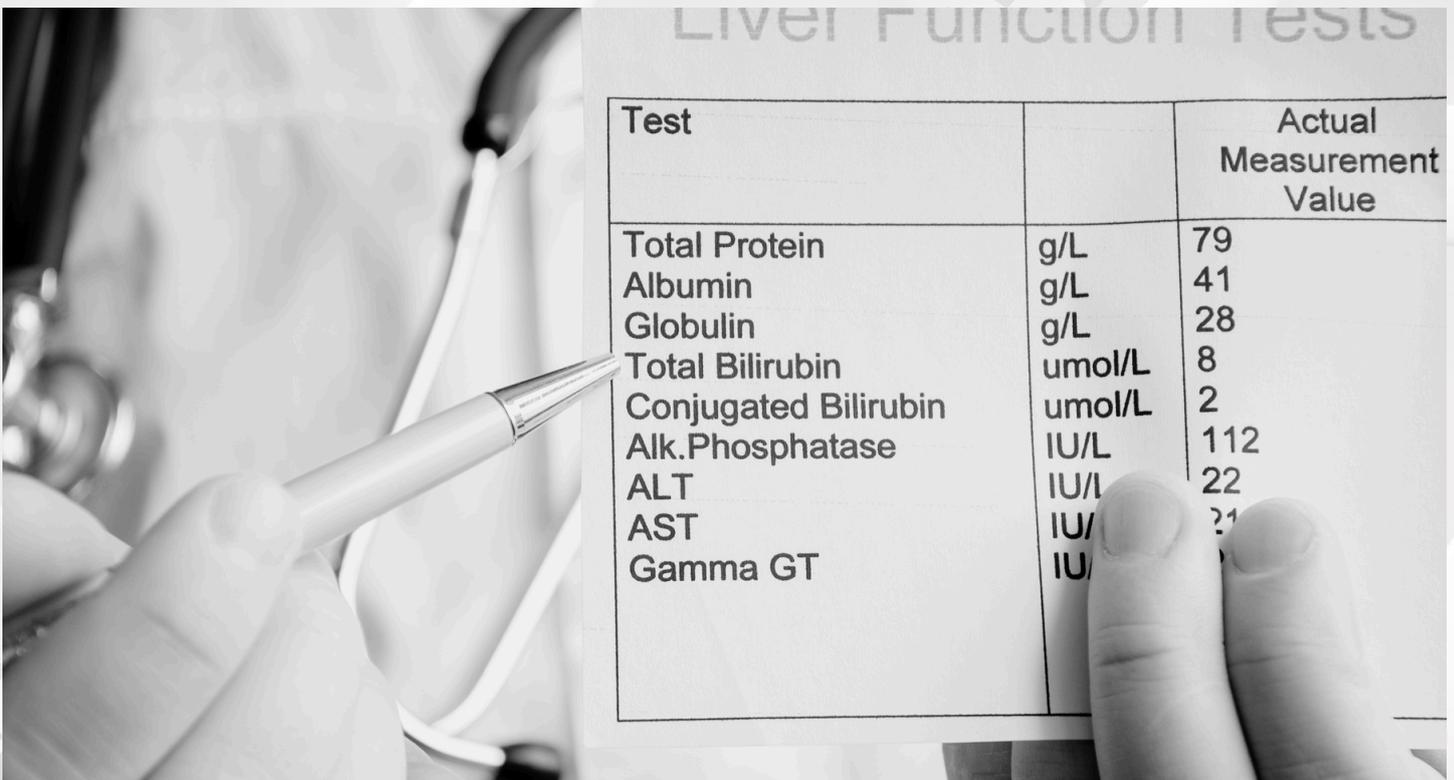
Data from **case reports** are examined for **peak values of liver enzymes ALT and TBL** over the observation period. These values are plotted on an x-y chart as logarithm10 multiples of elevations above the upper limits of the normal reference ranges (ULN).

# 2

For an individual patient, time course of ALT, TBL, AST, and ALP are plotted together for visual comparison.

# 3

A **medical text narrative, written by a skilled physician, provides additional context** about the patient's condition. This narrative helps estimate the most likely cause of abnormal findings and assesses the probability of drug-induced hepatotoxicity.

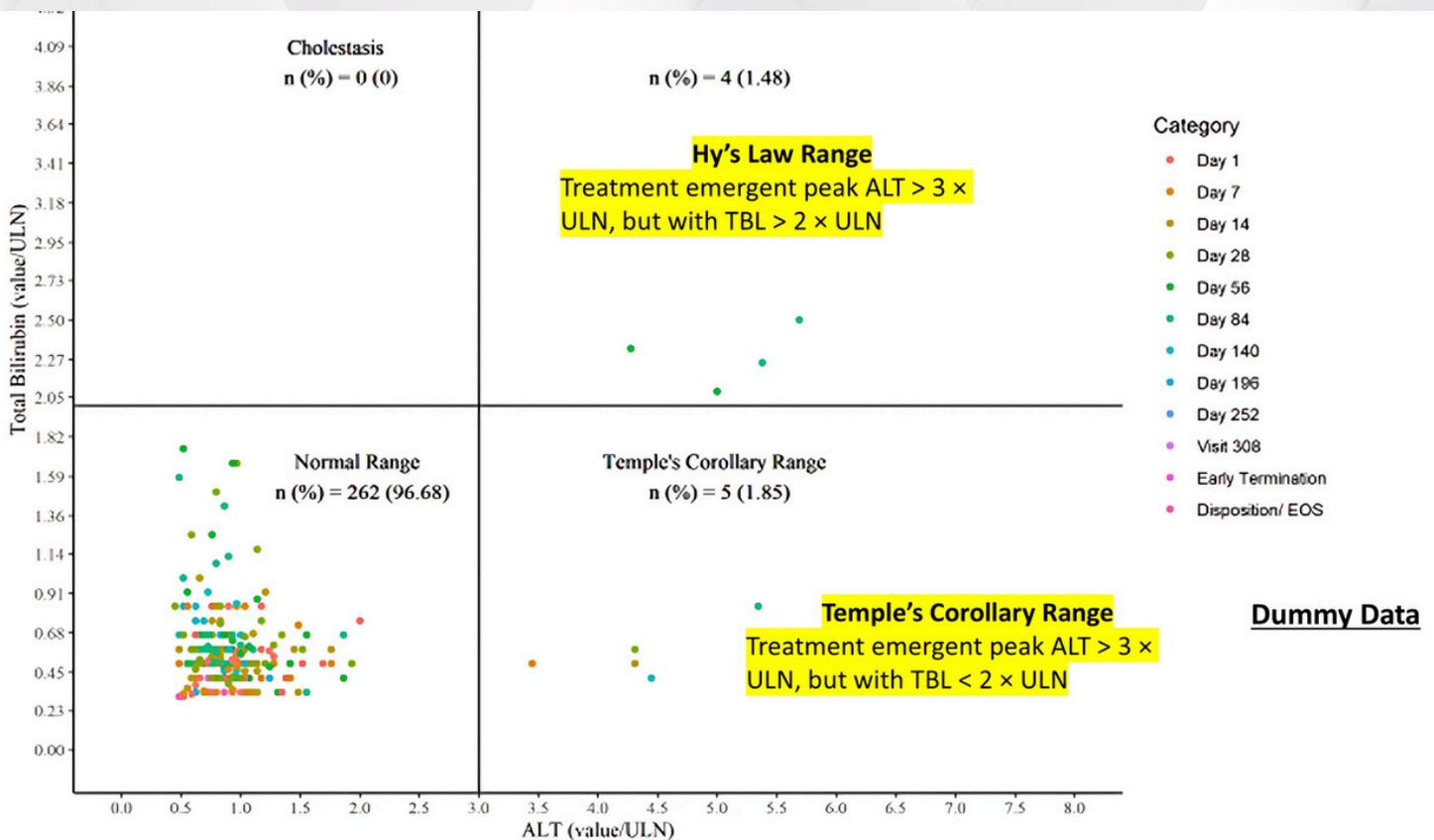


Test		Actual Measurement Value
Total Protein	g/L	79
Albumin	g/L	41
Globulin	g/L	28
Total Bilirubin	umol/L	8
Conjugated Bilirubin	umol/L	2
Alk.Phosphatase	IU/L	112
ALT	IU/L	22
AST	IU/L	21
Gamma GT	IU/L	21

# FDA's eDISH Program for Hepatotoxicity Assessment

## Causality Assessment :

- Requires considering of multiple potential factors.
- Several possible causes are common and Insufficient to simply label cases as 'confounded.'
- Estimated likelihood is categorized as 'probable' if the likelihood is over 50% and higher than all other causes combined.
- Sufficient information and thorough patient investigation are essential to rule out alternative causal factors.



**Note:** *The upper right quadrant doesn't automatically define cases as 'Hy's Law'; It identifies patients as of **unique importance**.* Further clinical information is essential for a comprehensive medical diagnosis aimed at identifying the most likely cause of the observed findings.

# Approach to the diagnosis of DILI

Abnormal biochemistry/acute hepatitis

Suspicion of DILI

## Features indicating toxic etiology

- Skin involvement
- Kidney injury
- Previous DILI episodes

Careful enquiry of exposure to Drugs, HDS, OTC (record start and stop dates)

## Potential drawbacks

- Lack of information (e.g., dose, duration)
- Various medications
- Hidden OTC and HDS intake
- Discontinue any non-essential drugs/HDS Treatments
- Search in hepatotoxicity resources (Liver tox)

Calculate biochemical pattern of liver injury

### Cholestatic

$$R = (\text{ALT/ULN}) / (\text{ALP/ULN}) \leq 2$$

### Mixed Injury

$$R > 2 \text{ or } < 5$$

### Hepatocellular

$$R = (\text{ALT/ULN}) / (\text{ALP/ULN}) \geq 5$$

Look for alternative causes

- Viral infections (HAV, HBV, HCV, HEV, EBV, CMV)
- Alcohol-related liver disease
- Hepatic ischaemia
- Autoantibody titres,  $\uparrow$  IgG
- Benign/malignant biliary obstruction
- Primary biliary cholangitis
- Primary sclerosing cholangitis

Liver biopsy maybe considered if

- Negative or incomplete dechallenge
- Acute or chronic atypical presentation
- Hepatic vascular disorder (e.g. ascites)
- Chronic hepatitis fibrosis
- Microvesicular steatosis
- Autoimmune hepatitis



# Conclusion

DILI is a **key concern** for regulators, drug developers, and physicians, and is **difficult to predict during drug development process.** 01

As severe DILI is generally rare, finding a single case may require treatment of thousands of people from varied patient populations. 02

The clinical trials present an exclusive opportunity to detect hepatotoxicity and cases of potential DILI with a study drug prior to its use in general population. 03

**Monitoring the liver test** abnormalities is useful for assessing trends over time and to **analyze imbalance between study drug and placebo/comparator groups.** 04

Due to the **limited number of subjects in a clinical trial, monitoring the standard serum liver tests to detect milder liver injury can be considered a predominant approach to predict the risk of possible DILI in clinical trials.** 05

Considering that there may be varied mechanisms of DILI and different clinicopathological phenotypes, a systematic collection of adequate diagnostic datasets along with a focused causality assessment performed by clinical experts is required for evaluation of each potential case of DILI in clinical trials. 06



# Some of the Key References

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# About Author

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## DR. SUMIT VERMA

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*Dr. Sumit Verma is a medical graduate with specialization in anesthesiology and has more than 15 years of experience in the pharmaceutical industry, clinical medicine, clinical research, and pharmacovigilance. He has built teams that have consistently delivered and exceeded customer expectations across pharmacovigilance domains such as case processing, signal management, risk management, aggregate reports, and clinical safety. He has co-authored two books – one on pharmacovigilance and another on pharmacology.*

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