EPA United States Environmental Protection Agency

Cross-Laboratory Comparison of Non-Targeted Analysis Method Performance using Complex Synthetic Mixtures

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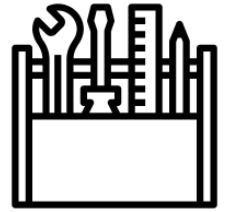
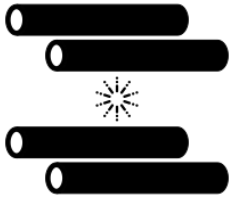
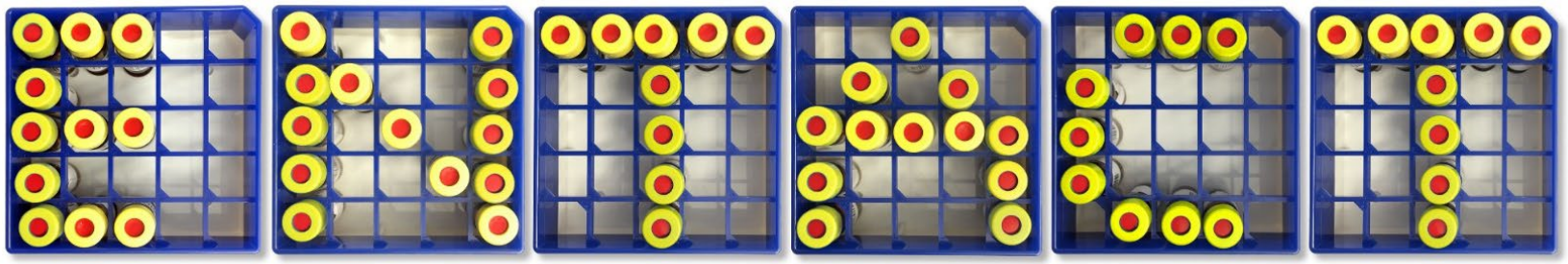
Science Questions for Research Community

- How variable are tools and results from lab to lab?
- Are some methods/workflows better than others?
- How does sample complexity affect performance?
- What chemical space does a given method cover?







EPA's Non-Targeted Analysis Collaborative Trial

EPA United States Environmental Protection Agency

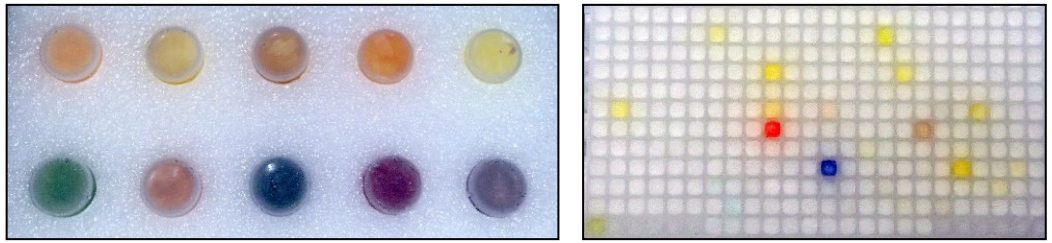
ENTACT Part 1 ★

Chemicals from ToxCast Library

~1200 ToxCast Chemicals (highest quality)

10 Mixtures (100-400 chemicals each)

*Multi-Well Plates**



~25 Collaborators & 6 Contractors*:

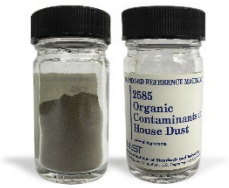
1st: Blinded analysis

2nd: Unveiling of chemicals

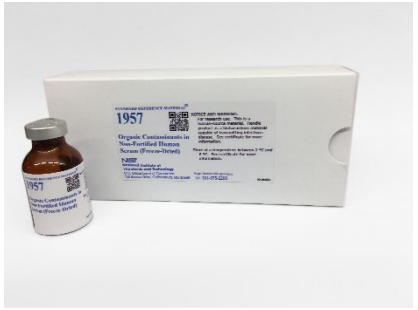
3rd: Unblinded evaluation

ENTACT Part 2

Reference & Fortified House Dust

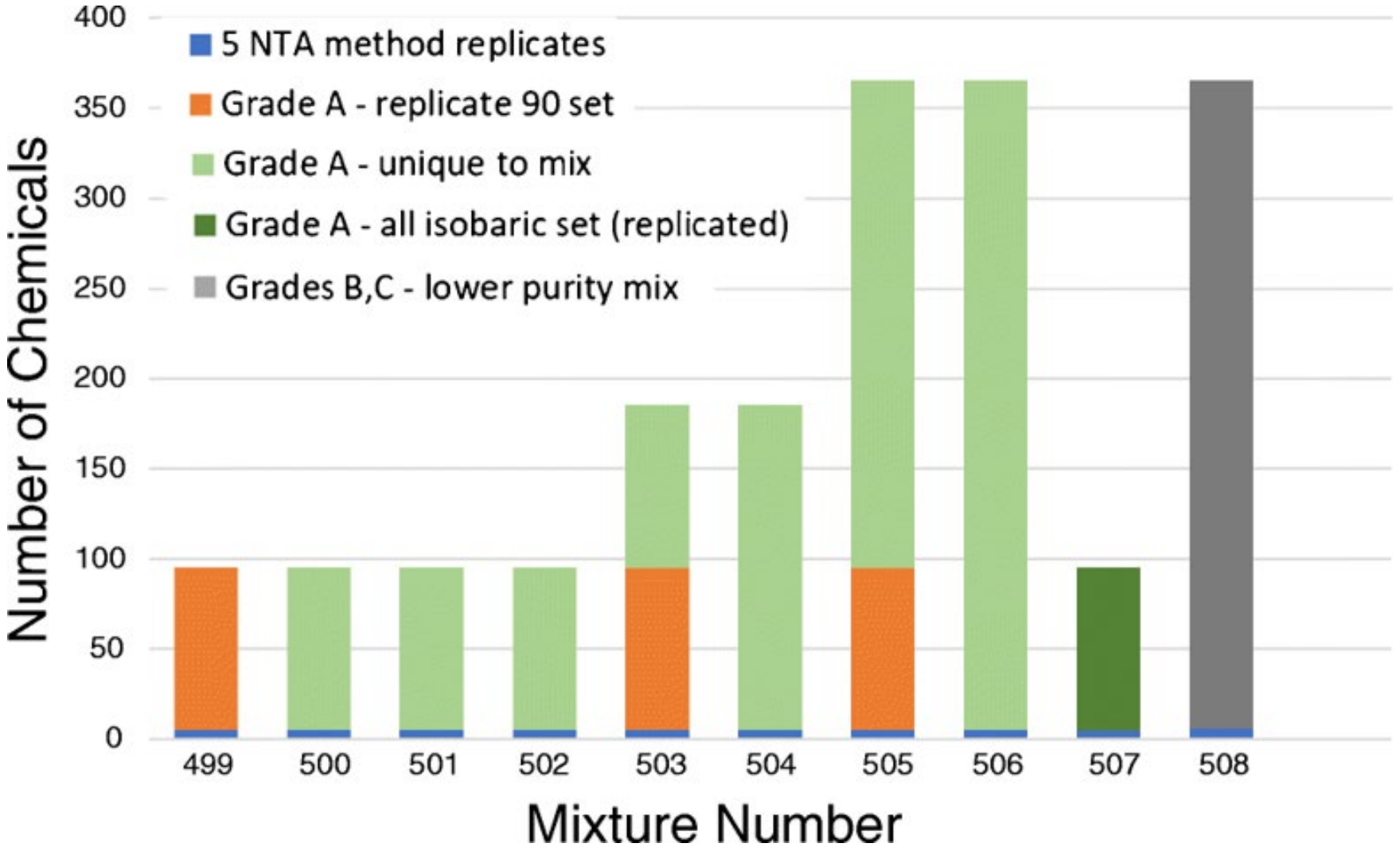


Reference & Fortified Human Serum



Reference & Fortified Silicone Wristbands

Design of ENTACT Mixtures



Replication in substance spikes offers a unique means to assess NTA method reproducibility!

Ulrich et al. 2019. doi: 10.1007/s00216-018-1435-6

Who is Working on ENTACT?

Contractors:

University of Alberta
Emory University
Duke University
Oregon State University OSU
San Diego State University
Pacific Northwest National Laboratory

19 blinded analysis submissions, to date…

Vendors:

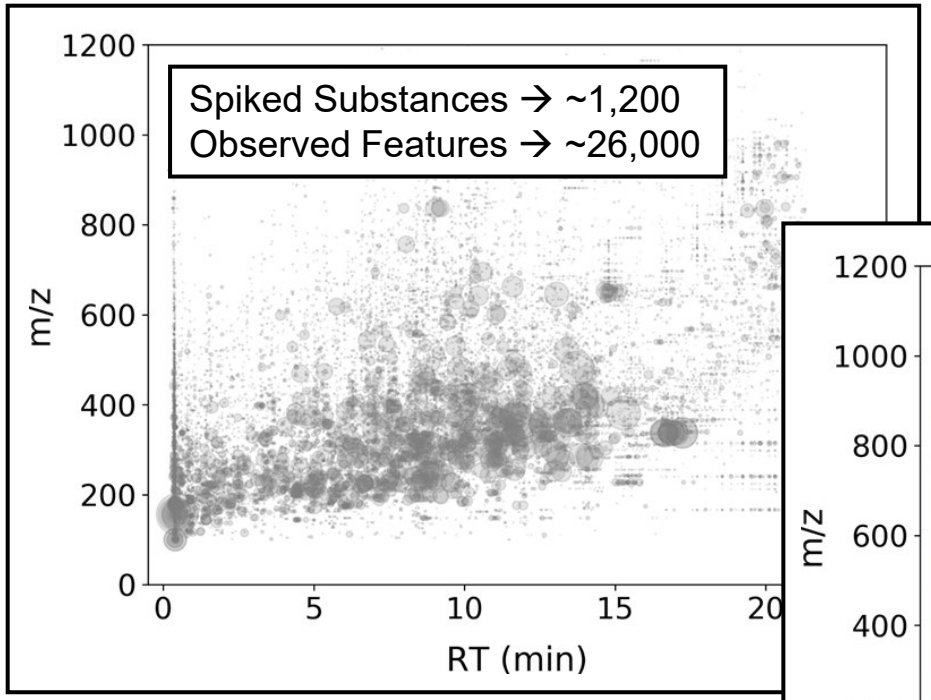
The Scripps Research Institute
Agilent Technologies
Thermo Scientific
Leco
AB SCIEX
Waters The Science of What's Possible.

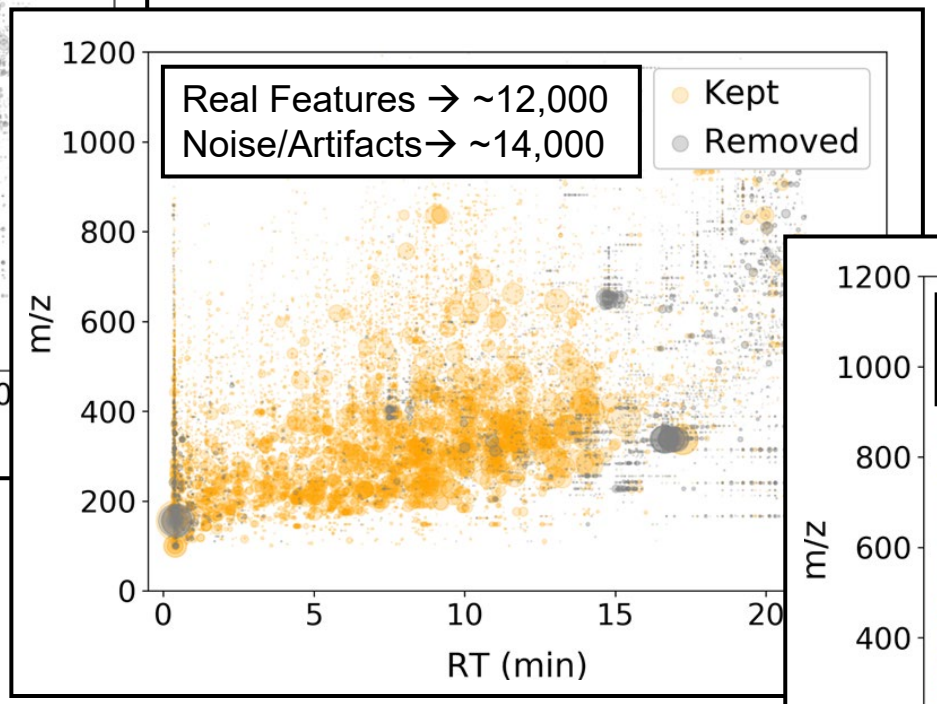
General Participants:

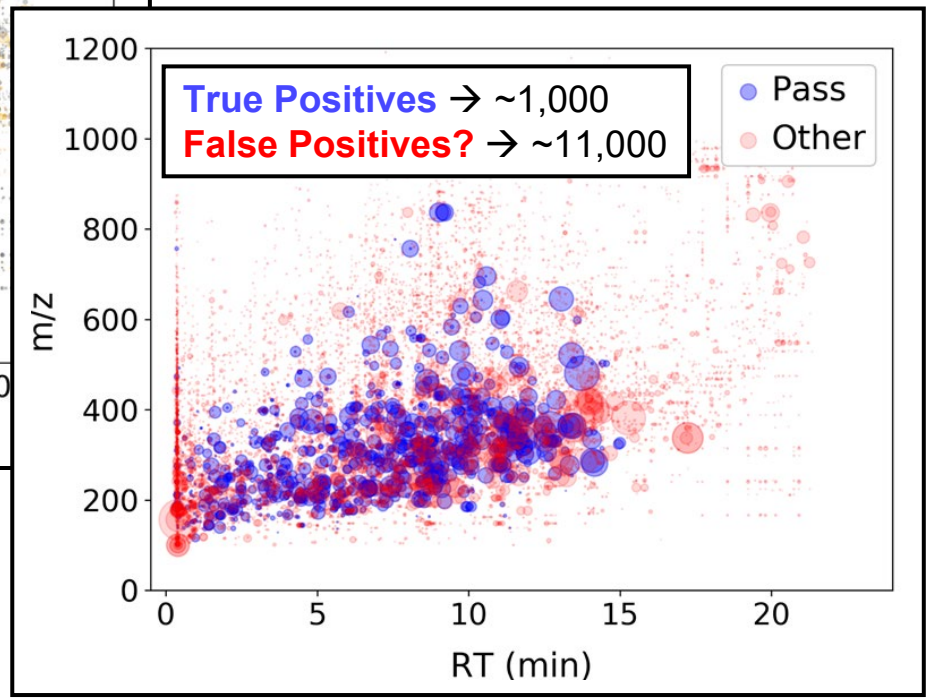
uni.lu Université du Luxembourg
UF University of Florida
Wisconsin State Laboratory of Hygiene, University of Wisconsin-Madison
NIST
NC State University
Recetox
Cornell University Founded A.D. 1865
UW Tacoma
Mount Sinai
University of Birmingham
DTSC Department of Toxic Substances Control
eawag aquatic research
United States Environmental Protection Agency
FIU

EPA Lab Results for ENTACT Mixtures

LC-QTOF HRMS (ESI+ and ESI-)



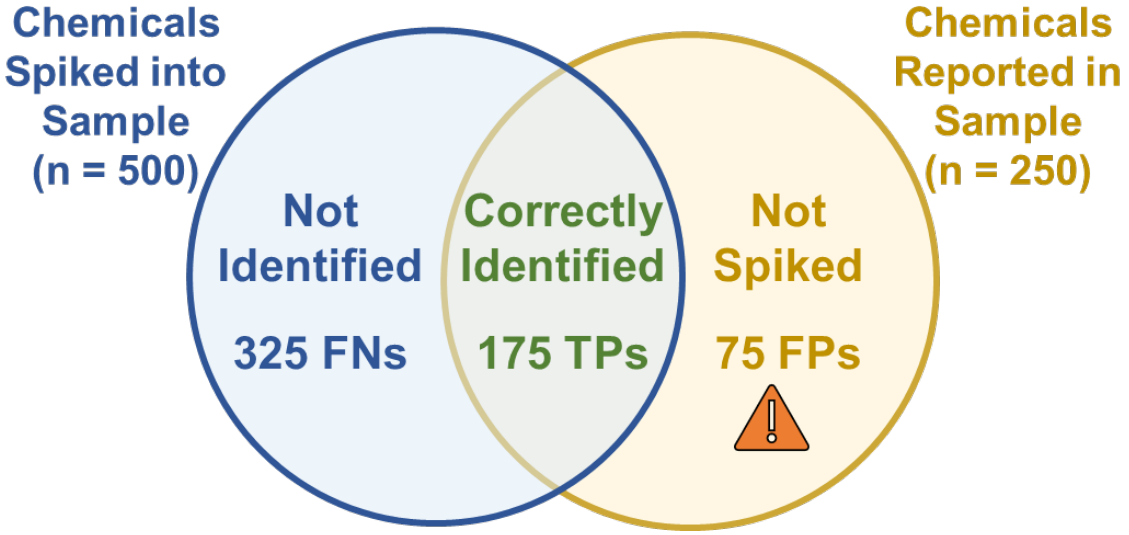




Substance Identified?	Substance Spiked? Yes	Substance Spiked? No
Yes	True Positives (≤ 65%)	False Positives?
No	False Negatives (≥ 35%)	True Negatives?

Sobus et al. 2019. doi: 10.1007/s00216-018-1526-4

Evaluation Tools Must Be Used With Caution



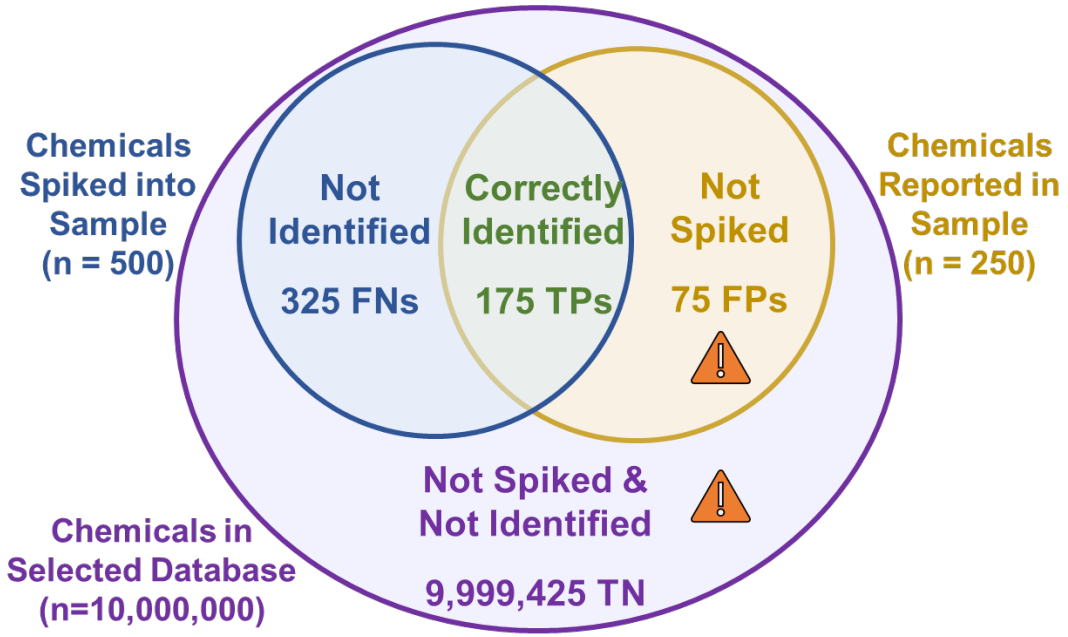
Boundary n = 575		Chemical is… spiked into sample	Chemical is… not spiked into sample		
Chemical is…	reported in sample	TP 175	FP 75 ⚠	Precision 0.70 ⚠	FDR 0.30
	not reported in sample	FN 325			
		TPR 0.35			
		FNR 0.65			

A hypothetical example

Fisher et al. 2022. doi: 10.1007/s00216-022-04203-3

- How do we differentiate FPs from unintentional TPs?
- What about lower-level “hits”? (e.g., Level 4)
- What about ID reproducibility?

Evaluation Tools Must Be Used With Caution



Boundary n = 10,000,000		Chemical is… spiked into sample	Chemical is… not spiked into sample		
Chemical is…	reported in sample	TP 175	FP 75 ⚠	Precision 0.70 ⚠	FDR 0.30
	not reported in sample	FN 325	TN 9,999,425 ⚠		
		TPR 0.35	FPR 0.00001 ⚠	F_1 0.47 ⚠	Accuracy 0.99996
		FNR 0.65	TNR 0.99999	MCC 0.49	

Fisher et al. 2022. doi: 10.1007/s00216-022-04203-3

- How do we appropriately handle TNs?

Processing ENTACT Data Submissions

- Individual methods treated separately (if appropriate)
- One candidate mass/formula/compound per feature
- Confidence level revised as needed (with consensus)
- Matching to spiked substances by mass, formula & structure
- "**Observed**" if structure or formula (no spiked isomers) match
- "**Correctly Identified**" if structure match
- "**Reproducible**" if correctly ID'd >50% of the time
  - "Eligible" compounds spiked >1 time and identified ≥1 time

Forward vs. Reverse Evaluation

Forward

Spiked Compound	Observed?	Correctly ID'd	Reproducibly ID'd
1	Yes	Yes	Yes
2	Yes	Yes	No
3	No	No	--
4	No	No	--
5	Yes	Yes	Yes
6	Yes	No	--
7	Yes	Yes	No
8	No	No	--
9	Yes	Yes	--
10	No	No	--
…	…	…	…
100	Yes	Yes	Yes

Reverse

Reported Compound	Spiked?
1	No
2	Yes
3	Yes
4	Yes
5	No
6	No
7	Yes
8	No
9	Yes
10	Yes
…	…
125	No

Outlining Utilized Performance Metrics

- **Observability Rate** = # Observed / # Spiked
  - If it was spiked, could your instrument detect it?
- **True Positive Rate** = # Correctly Identified / # Spiked
  - If it was spiked, could your workflow correctly ID it?
- **Correct ID Rate** = # Correctly Identified / # Observed
  - If it was observed, could your workflow correctly ID it?
- **Reproducibility Rate** = # Reproducible / # Eligible
  - If it was correctly ID'd once, was it correctly ID'd most of the time?

Forward Evaluation Metrics

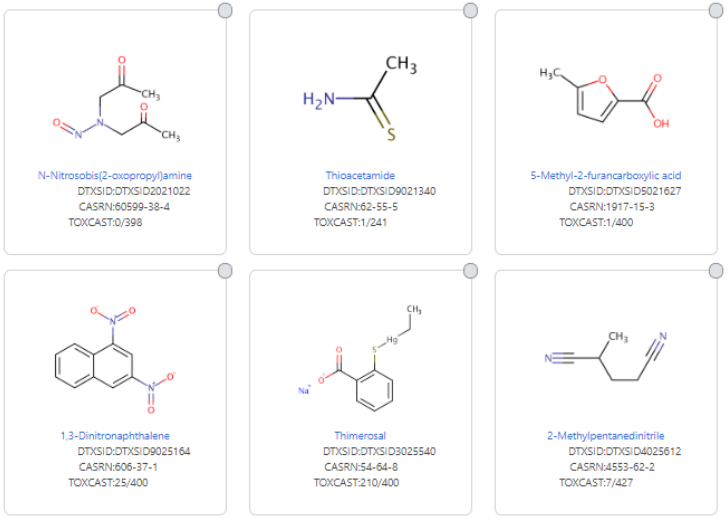
- **Reporting Rate** = # Reported / # Spiked
  - What is the ratio of reported to spiked compounds?
- **Correct Reporting Rate** = # Correctly Identified / # Reported
  - If it was reported, was it a correctly identified spiked compound?

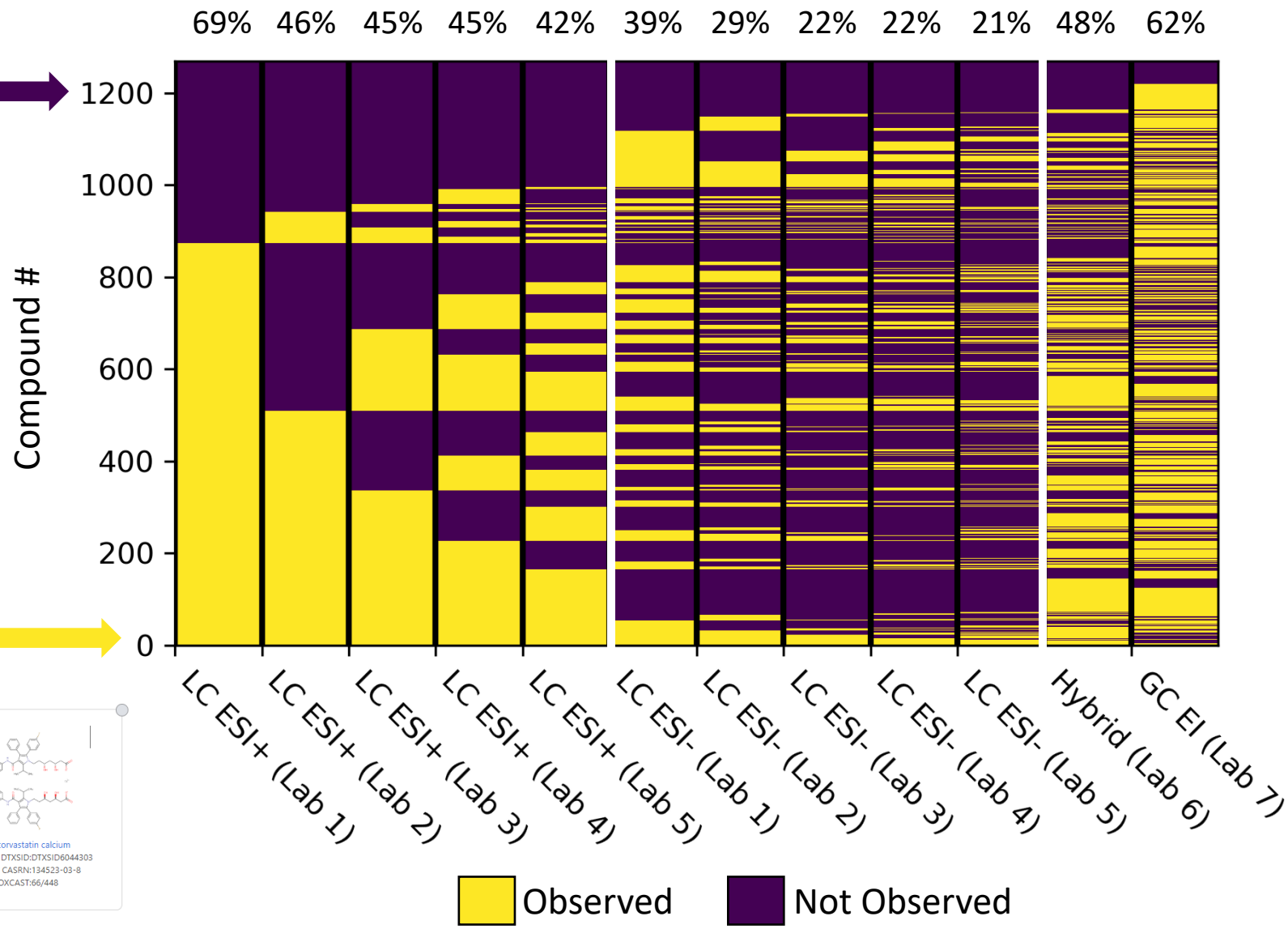
Reverse Evaluation Metrics

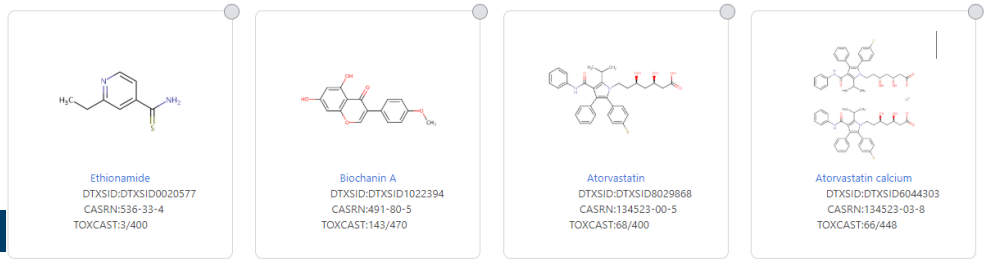
Method Comparison: “Observed” Compounds

7 Labs, 12 Methods

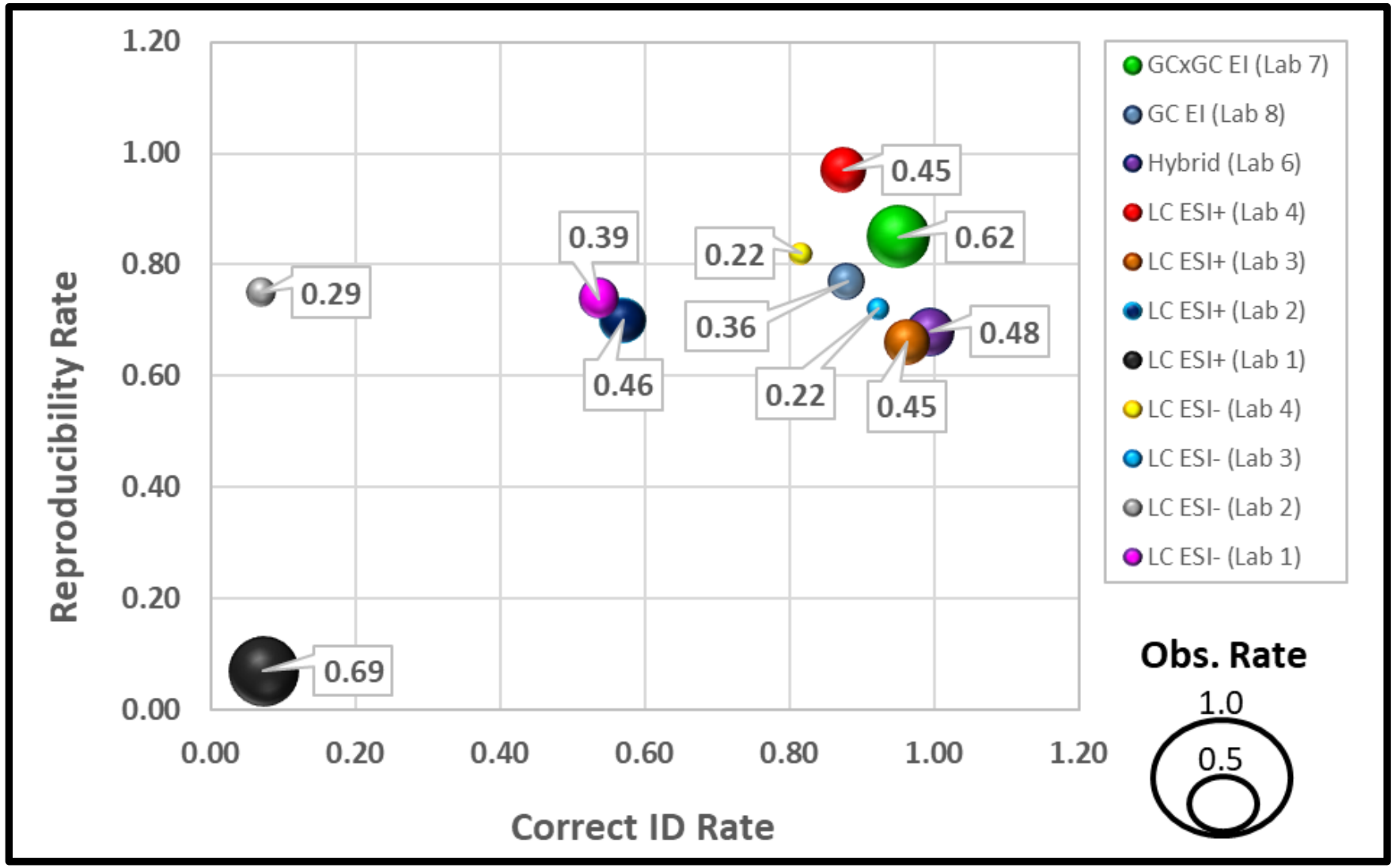
~5% Not Observed by Any Method







Method Comparison: 3 Forward Metrics



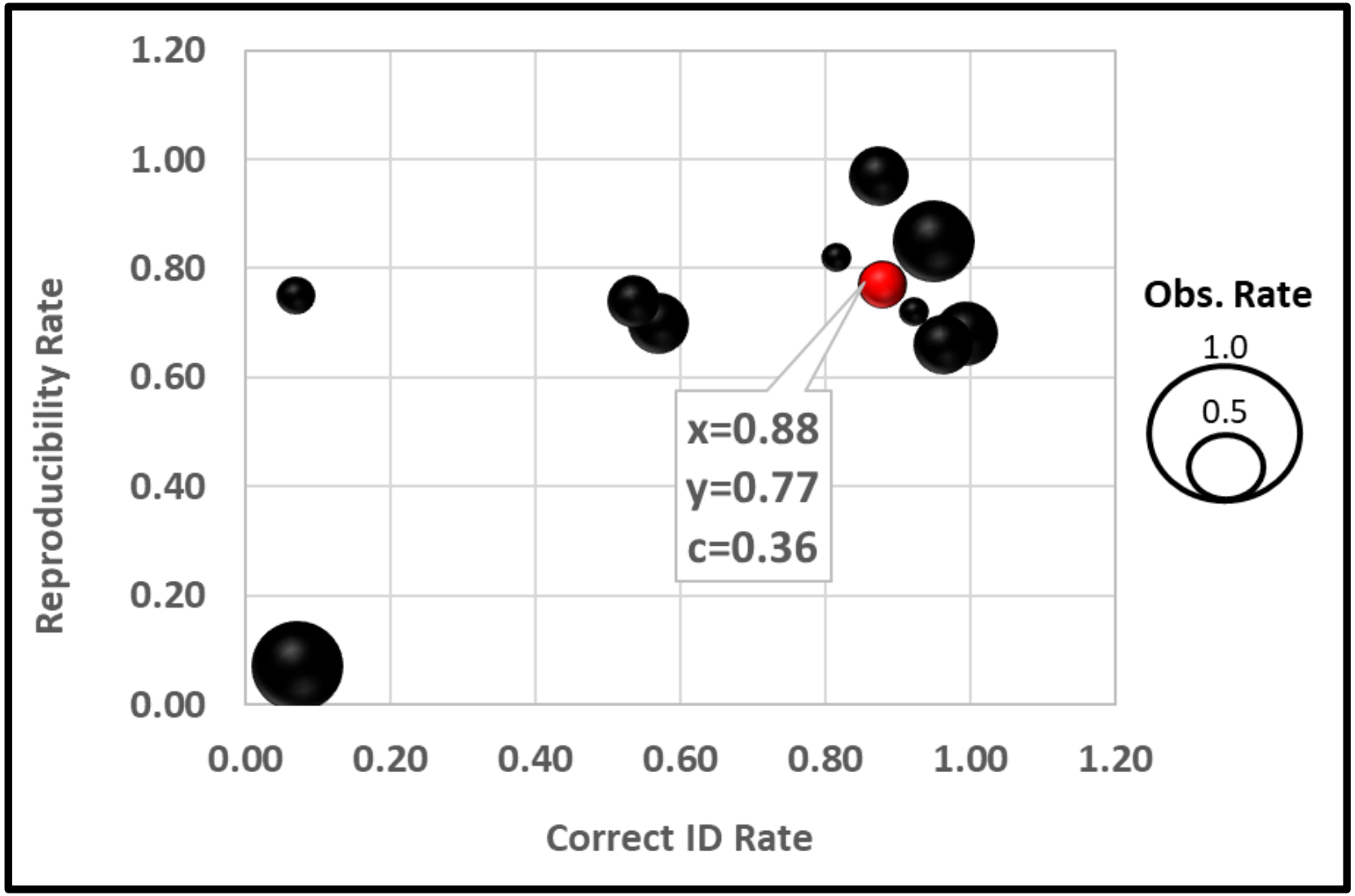
Metrics (all %):

Bubble Size →
How often observed?

X-Axis →
How often correctly ID'd if observed?

Y-Axis →
How consistently ID'd?

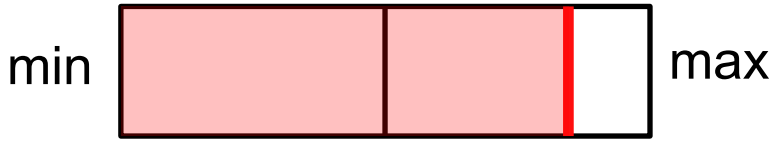
Example Performance Report



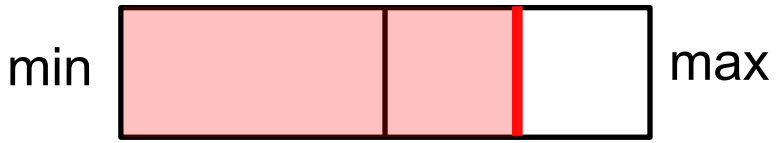
Performance Scores:

(% of max score)

Correct ID Rate: **88%**

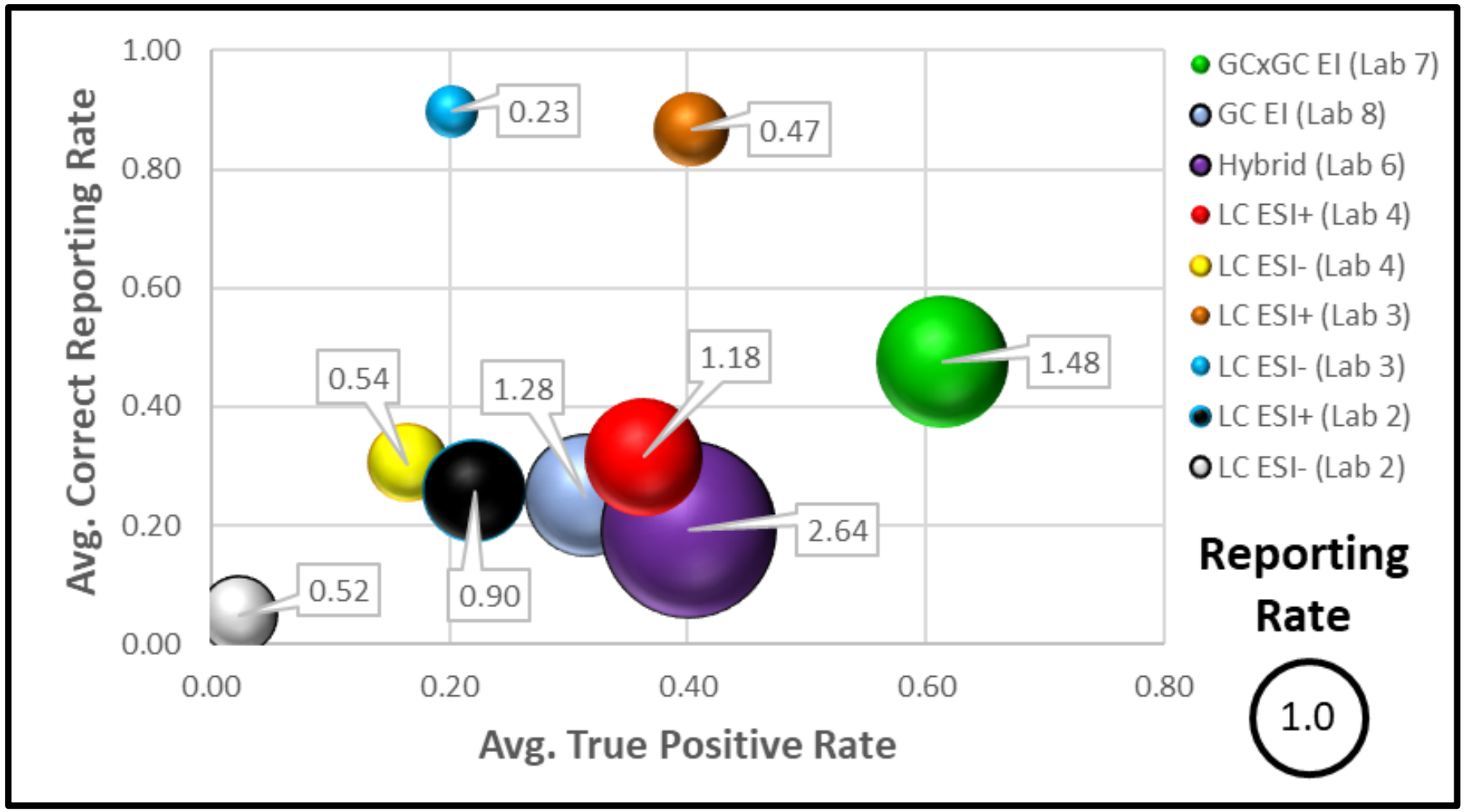


Repro. Rate: **78%**



Obs. Rate: **30%**

Method Comparison: TPR & Reverse Metrics



Metrics (all %):

Bubble Size → Amount reported

X-Axis → Correct IDs across all spikes

Y-Axis → Correct IDs across all reported

Additional Results for Collaborators

- Simple performance summary file (n=1 per method):
  - # and % correct identifications per sample
- Individual results files (n=10 per method):
  - Mass match (yes/no), formula match (yes/no), compound match (yes/no)
  - Highest confidence level (as reported or after consensus revision)
- Composite results file (n=1 per method):
  - For each spiked substance (n=1,269)
    - # of spikes (1-10), # of isomer spikes (1-5)
    - # mass hits, # formula hits, # compound hits
    - Observed (yes/no/undetermined), Correct ID (yes/no), Reproducible (yes/no)

Some Challenges (to date)

- Multiple chemical candidate submissions per feature
- Inconsistent & inaccurate use of scoring metrics
- Inconsistent & inaccurate chemical reporting procedures
- Inconsistent and unclear feature filtering protocols
- Limited engagement regarding collaborator follow-up
- Determining FPs vs. uTPs
- Determining TNs and dependent metrics
- Slow evaluation process vs. rapid method development processes

Summary of ENTACT Findings

- **NTA methods are suitable for many ToxCast chemicals**
  - ~5% of ENTACT compounds not observed by any method
- **Multiple methods required for broad characterization**
  - No “one size fits all” method
  - <1% of ENTACT compounds observed using all methods
- **Performance determined across multiple metrics:**
  - **Observability Rate** = Ability to observe those spiked → (22% to 69%)
  - **True Positive Rate** = Ability to identify those spiked → (2% to 61%)
  - **Correct ID Rate** = Ability to identify those observed → (7% to 99%)
  - **Reproducibility Rate** = Ability to consistently identify → (7% to 97%)
  - **Reporting Rate** = Amount reported vs. spiked → (23% to 264%)
  - **Correct Reporting Rate** = Amount correctly ID’d vs. reported → (5% to 90%)

Questions?

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