

1           **Systematic bias in network proximity Z-scores: A**  
2           **comparative robustness audit using *Hypericum perforatum***  
3           **constituents**

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5           **Abstract**

6     Network-based metrics are widely used to identify associations between compounds and diseases, as-  
7     suming that proximity within a protein–protein interaction network reflects functional relevance. How-  
8     ever, these metrics are often reported as Z-scores, which we demonstrate are fundamentally sensitive to  
9     the number of targets a compound possesses. This dependency introduces a systematic bias where com-  
10    pounds with broad polypharmacology appear statistically significant due to null distribution tightening  
11    (the Law of Large Numbers) rather than physical network reachability. Here, we systematically evaluate  
12    this bias using the human liver interactome and a controlled comparison of two constituents from *Hy-*  
13    *pericum perforatum*. We show that conventional proximity Z-scores yield unstable rankings that reverse  
14    depending on network construction parameters. While a compound with many targets may achieve a  
15    higher Z-score, it can remain physically more distant from the disease module than a compound with  
16    fewer, high-leverage targets. We resolve this by utilizing random walk–based influence propagation and  
17    introducing a size-normalized metric: perturbation efficiency. Our results show that influence-based  
18    rankings are stable across varied network thresholds and correctly identify high-leverage modulators that  
19    proximity metrics miss. This study provides a methodological template for identifying and correcting  
20    statistical artifacts in network medicine, enabling more reliable risk assessment in complex biological  
21    systems.

22    **Keywords:** network propagation, proximity metrics, metric robustness, drug-induced liver injury,  
23    polypharmacology, Z-score confounding, perturbation efficiency.

## 24 1 Introduction

25 Network-based prioritization is a cornerstone of modern systems biology and drug discovery, assuming  
26 that the topological proximity between compound targets and disease genes within a protein–protein in-  
27 teraction (PPI) network reflects functional relevance [1–4]. Because raw network distances are sensitive  
28 to local topology and degree distribution, they are typically reported as Z-scores relative to degree-  
29 matched null models. While these Z-scores successfully quantify statistical significance in most applica-  
30 tions [3], we demonstrate they can be confounded by large asymmetries in target set size. As the number  
31 of seed nodes increases, the variance of the null distribution decreases (the Law of Large Numbers),  
32 leading to deterministic significance inflation for compounds with broad polypharmacology. Identifying  
33 whether such results reflect true biological influence, or whether they represent systematic artifacts of  
34 distance-based inference, is essential for the reliability of network medicine.

35 Using the human liver interactome as a model system, we investigate this confounding effect through  
36 a controlled comparison of two constituents from *Hypericum perforatum* (St. John’s Wort). These con-  
37 stituents—Hyperforin and Quercetin—exhibit highly asymmetric target set sizes: Hyperforin possesses  
38 10 validated targets, while Quercetin has over 60 [5–7]. This asymmetry provides a sharp stress test  
39 for network metrics. Conventional proximity Z-scores predict greater disease-associated significance for  
40 the broad-spectrum modulator, even when it is physically more distant from the disease module than the  
41 high-leverage modulator. This reversal indicates that proximity-based prioritization is unstable across  
42 network construction parameters and susceptible to sample-size artifacts.

43 Here, we evaluate the robustness of proximity-based and influence-based metrics for comparative  
44 prioritization. We demonstrate that proximity Z-scores yield unstable, threshold-dependent rankings  
45 driven by null-distribution tightening rather than physical reachability. To resolve this instability, we  
46 utilize random walk–based influence propagation, which integrates over the entire network topology and  
47 captures signal amplification through regulatory hubs [8]. We introduce a normalized metric, perturba-  
48 tion efficiency, to account for target set size and ensure unbiased comparisons. Our results show that  
49 influence-based propagation provides a stable, theoretically consistent framework for network pharma-  
50 cology that correctly identifies high-leverage perturbations where traditional proximity metrics fail.

## 51 2 Results

### 52 2.1 Proximity Z-scores are confounded by target set size

53 We first established network context by quantifying target count and shortest-path proximity to 82 DILI-  
54 associated genes (Figure 1). Quercetin engages 62 targets in the liver-expressed largest connected  
55 component; Hyperforin engages 10. At STRING confidence  $\geq 900$ , Hyperforin targets are physically  
56 closer to DILI genes ( $d_c = 1.30$ ) than Quercetin targets ( $d_c = 1.68$ ; Table 1). However, the proxim-  
57 ity Z-scores yield the opposite ranking: Quercetin achieves  $Z = -5.44$  ( $p < 0.001$ ), while Hyperforin  
58 achieves  $Z = -3.86$  ( $p < 0.001$ ). All reported associations survived Benjamini–Hochberg FDR correc-  
59 tion ( $q < 0.05$ ).

60 This discrepancy highlights a fundamental confounder in proximity Z-scores: the law of large num-  
61 bers. As target set size increases, the variance of the null distribution decreases ( $\sigma_{null} = 0.09$  for  
62 Quercetin vs 0.24 for Hyperforin), inflating the significance of broader target sets despite greater physical  
63 distance. This statistical artifact suggests that Quercetin poses greater risk, whereas the physical topology  
64 favors Hyperforin. This effect occurs independently of network construction parameters and represents  
65 a fundamental statistical property of averaged distributions: larger samples inherently produce more pre-  
66 cise (narrower) null distributions, artificially inflating Z-score magnitude regardless of actual topological  
67 proximity.

### 68 2.2 Influence-based rankings are stable and resolve the confound

69 Random walk with restart (RWR) stabilizes this ranking by integrating over all paths (Figure 2). Hyper-  
70 forin achieves influence  $Z = +10.12$  ( $p < 0.001$ ); Quercetin achieves  $Z = +4.55$  ( $p < 0.001$ ; Table 1).  
71 Unlike proximity, influence Z-scores correctly reflect the topological advantage of Hyperforin’s regu-  
72 latory hub occupancy. This ranking aligns with the known biological ground truth: Hyperforin is the  
73 hepatotoxic constituent responsible for drug-drug interactions via PXR activation, while Quercetin has  
74 no documented hepatotoxicity and may be hepatoprotective. The ranking remains consistent across  
75 topology-only and expression-weighted analyses, demonstrating that influence propagation is less sus-  
76 ceptible to sample-size artifacts than shortest-path distance.

### 77 2.3 Expression weighting refines the signal

78 To assess whether the RWR signal persists under tissue-specific constraint, we applied expression-  
79 weighted influence propagation (EWI), weighting transitions by destination-node liver expression (Fig-

80 ure 3).

81 The Z-score differential narrows but remains substantial under expression weighting: Hyperforin  
82  $Z = +8.98$  ( $p < 0.001$ ); Quercetin  $Z = +5.79$  ( $p < 0.001$ ). Hyperforin's advantage is driven primarily  
83 by the PXR–CYP master regulatory axis, which remains highly active in liver tissue (e.g., CYP3A4 at  
84 335 TPM). Quercetin's influence is moderated by its broad, diffuse target profile, which includes several  
85 high-expression nodes (e.g., CFB at 1,115 TPM) that do not converge on a DILI effector hub.

86 **2.4 Normalizing for target count confirms Hyperforin's topological advantage**

87 To resolve the target-count paradox, we compared the average network influence of each individual  
88 target, reframing polypharmacology as an efficiency problem rather than a coverage problem (Figure 4;  
89 Table 2).

Compound	Targets	Eff. (RWR)	Eff. (EWI)	RWR Ratio*	EWI Ratio†
Hyperforin	10	0.1138	0.1330	—	—
Quercetin	62	0.0322	0.0493	—	—
<b>Fold difference</b>	—	—	—	<b>3.5× (3.7×)</b>	<b>2.7× (2.8×)</b>

91 \*RWR Ratio: observed ratio (robust ratio in parentheses). †EWI Ratio: observed ratio (robust ratio in paren-  
92 theses).

93 Each Hyperforin target contributes  $3.7\times$  more DILI-directed influence than each Quercetin target (robust ra-  
94 tio). This disparity indicates that Hyperforin's target positions are substantially higher leverage than those of  
95 Quercetin, achieving greater perturbation efficiency despite a 6-fold smaller target set. Critically, this 3.7-fold  
96 efficiency advantage is threshold-independent: it holds at STRING  $\geq 700$  ( $3.8\times$ ),  $\geq 900$  ( $3.7\times$ ), and under both  
97 topology-only (RWR) and expression-weighted (EWI) regimes. Unlike Z-scores, which reverse between thresh-  
98 olds, perturbation efficiency provides a stable, parameter-invariant comparative metric.

99 **2.5 Bootstrap resampling excludes target-selection bias**

100 To rule out the possibility that Hyperforin's advantage arises from favorable target selection rather than strategic  
101 network positioning, we performed bootstrap sensitivity analysis (Figure 5). 100 random 10-target subsets were  
102 sampled without replacement from Quercetin's 62-target pool and scored by RWR.

103 Hyperforin's observed influence (0.1138) exceeds the entire bootstrap distribution from Quercetin (mean =  
104 0.0308, 95% CI = [0.0160, 0.0542]; Table 3). The fold difference between Hyperforin and the bootstrap mean is  
105  $3.7\times$ . This confirms that Hyperforin's advantage is not an artifact of target count; even when sampling equalized  
106 subsets from Quercetin's pool, no configuration matches Hyperforin's influence.

107 **2.6 Ranking stability across network thresholds**

108 The influence ranking is stable across network confidence thresholds (Table 6). Hyperforin ranks first in all RWR  
109 and EWI configurations at both  $\geq 700$  and  $\geq 900$  thresholds. Notably, the proximity ranking reverses between  
110 thresholds: at  $\geq 700$ , Hyperforin is physically closer ( $d_c = 0.60$  vs 1.34) and more "significant" ( $Z = -6.04$  vs  
111  $-5.46$ ). At  $\geq 900$ , Quercetin appears more "significant" ( $Z = -5.44$  vs  $-3.86$ ) despite being physically more dis-  
112 tant (1.68 vs 1.30). This instability in proximity Z-scores—while influence rankings remain stable—demonstrates  
113 that influence-based metrics are more robust to network construction parameters.

114 **2.7 Chemical similarity excludes structural confounding**

115 To exclude the possibility that Hyperforin's network signal reflects structural similarity to known hepatotoxins,  
116 we performed chemical similarity analysis against the DILrank reference set (Figure 6). Morgan fingerprints  
117 (ECFP4) revealed that neither compound exceeds the 0.4 Tanimoto threshold for structural analog detection. No-  
118 tably, Quercetin exhibits higher structural similarity to DILI reference drugs yet lower network influence, reinforc-  
119 ing that the observed asymmetry is driven by network topology rather than chemical features.

120 **3 Discussion**

121 **3.1 Ranking stability and the Z-score confound**

122 The results of this study highlight a potential limitation in the use of network proximity Z-scores when comparing  
123 compounds with asymmetric target set sizes. While proximity is a standard prioritization criterion, our analysis  
124 demonstrates that its significance rankings can be influenced by the target count rather than topological distance  
125 alone. As the number of targets increases, the variance of the null distribution decreases (a manifestation of the Law  
126 of Large Numbers), which can lead to inflated significance levels for compounds with broad polypharmacology. In  
127 our controlled comparison, this effect causes a reversal of proximity-based rankings between network thresholds,  
128 failing to accurately reflect the physical distance advantage of a high-leverage modulator.

129 Influence-based metrics (RWR and EWI) appear less sensitive to this particular artifact. By integrating over the  
130 entire network topology, these methods capture signal propagation through regulatory hubs, providing rankings that  
131 remain stable across different network construction parameters. This relative stability suggests that influence-based  
132 propagation may offer a more robust framework for comparative network medicine, particularly in the presence of  
133 incomplete or asymmetric pharmacological data.

134 The mechanistic explanation for this robustness is that RWR integrates over *all* paths, capturing how signals  
135 amplify through hubs like PXR and AKT1. Shortest-path proximity, by contrast, is a descriptive metric for min-  
136 imum reachability; treating it as an inferential surrogate for functional impact conflates topological context with  
137 biological consequence.

<sup>138</sup> **3.2 Expression weighting as a biological constraint**

<sup>139</sup> Expression-weighted influence (EWI) constrains signal propagation to liver-active nodes. By attracting signal to  
<sup>140</sup> highly expressed proteins (destination-node weighting), we ensure that the network propagation reflects tissue-  
<sup>141</sup> specific biology. Under this constraint, the Hyperforin advantage persists, demonstrating that its topological ef-  
<sup>142</sup> ficiency is not an artifact of an unconstrained PPI network but is supported by the expression profile of the liver.  
<sup>143</sup> Attenuation of signal is expected when walks are constrained to active pathways; the fact that the ranking remains  
<sup>144</sup> stable provides positive evidence for the biological relevance of the PXR axis.

<sup>145</sup> **3.3 Perturbation efficiency vs. topological coverage**

<sup>146</sup> By normalizing total influence for target set size (where the restart vector is already  $|T|$ -weighted), we provide  
<sup>147</sup> a more balanced comparison of perturbation efficiency. Our results show that a single Hyperforin target exerts  
<sup>148</sup> 3.7-fold more influence on the DILI module than a Quercetin target.

<sup>149</sup> This efficiency claim is further validated by bootstrap sensitivity analysis. Even when sampling size-matched  
<sup>150</sup> 10-target subsets from Quercetin's pool, none reached the influence level achieved by Hyperforin. This demon-  
<sup>151</sup> strates that the advantage is not due to target count, but to the strategic network position of Hyperforin's tar-  
<sup>152</sup> gets—specifically their convergence on the PXR master regulator and downstream CYP effectors.

<sup>153</sup> **3.4 Mechanistic context: The PXR axis**

<sup>154</sup> The stability of the influence ranking aligns with the well-characterized PXR–CYP master regulatory axis. Hyper-  
<sup>155</sup> forin's primary target, NR1I2 (PXR), induces the expression of major xenobiotic metabolism enzymes including  
<sup>156</sup> CYP3A4 and CYP2C9 [6, 9]. In our network analysis, these effectors are part of the target set and the DILI  
<sup>157</sup> module, creating a high-connectivity hub structure that enables efficient propagation. Quercetin's 62 targets, while  
<sup>158</sup> numerous, are distributed across redundant or peripheral pathways that do not converge on a regulatory bottleneck.  
<sup>159</sup> Furthermore, clinical evidence indicates that Quercetin is not associated with hepatotoxicity and may exhibit hep-  
<sup>160</sup> atoprotective properties [7, 10]. Recent experimental studies have corroborated that St. John's wort exacerbates  
<sup>161</sup> hepatotoxicity through precisely this PXR-mediated bioactivation mechanism [11].

<sup>162</sup> **3.5 Limitations**

<sup>163</sup> Several limitations warrant consideration. First, network influence is a measure of topological reach and pertur-  
<sup>164</sup> bation potential, not a direct surrogate for toxicological outcomes. This model is dose-independent and does not  
<sup>165</sup> account for pharmacokinetics, binding affinity, or saturation kinetics. A high influence score indicates that a com-  
<sup>166</sup> pound's targets are strategically positioned to modulate a disease module, but the actual biological effect depends  
<sup>167</sup> on the molecular mechanism of action (e.g., agonism vs. antagonism) and the kinetic context.

<sup>168</sup> Second, while we demonstrate that proximity Z-scores are confounded by target set size, influence-based Z-  
<sup>169</sup> scores are not entirely immune to this effect. As the number of seed nodes increases, the variance of the null

170 distribution for influence sums also decreases, though less severely than for distance-based metrics. Critically,  
171 our core claims do not rest on absolute Z-score comparisons. We demonstrate that influence-based *rankings* are  
172 stable across network thresholds, while proximity rankings are not. We further resolve the size-dependence by  
173 introducing perturbation efficiency (influence per target), which explicitly normalizes for target count and provides  
174 a bias-corrected comparative metric.

175 Third, our case study is limited to a single botanical with two contrasting constituents. While this provides a  
176 controlled minimal model, generalization to larger compound libraries will require further validation.

### 177 **3.6 Conclusions**

178 In this study, we utilized *H. perforatum* as a known toxicological model to validate the *reliability* of network  
179 metrics; the biological ground truth (Hyperforin-mediated PXR activation) allowed us to confirm that influence  
180 propagation correctly identifies high-leverage perturbations where proximity metrics fail. The methodological  
181 conclusion is that proximity Z-scores are susceptible to sample-size confounding and should be used descriptively  
182 rather than for comparative inference across compounds with differing target counts. Influence-based propagation,  
183 combined with per-target normalization, provides a more stable framework that survives robustness checks and  
184 aligns better with mechanistic reality.

185 More broadly, this work provides a methodological template for identifying and resolving metric artifacts  
186 in network toxicology. By integrating signed edge weights and transcriptometric data, future iterations of this  
187 framework could investigate phenotype-specific associations, linking topological influence on specific biological  
188 sub-modules to discrete clinical outcomes.

## 189 **4 Methods**

### 190 **4.1 Data sources**

#### 191 **4.1.1 Protein–protein interaction network**

192 Human protein–protein interactions were obtained from STRING v12.0 [12]. Combined confidence scores were  
193 computed per STRING methodology (text mining, experiments, databases, co-expression, neighborhood, gene  
194 fusion, co-occurrence). Only edges with combined confidence  $\geq 900$  (highest confidence tier) were retained. Raw  
195 network: 11,693 genes, 100,383 edges.

#### 196 **4.1.2 Liver expression data**

197 Gene expression data were obtained from the Genotype-Tissue Expression Project (GTEx) v8 [13]. Median tran-  
198 scripts per million (TPM) values for liver tissue were extracted from the 2017-06-05 release (RNASeQCv1.1.9).  
199 Genes with liver TPM  $\geq 1$  were retained. Result: 13,496 liver-expressed genes.

200 **4.1.3 Drug-induced liver injury gene set**

201 DILI-associated genes were obtained from DisGeNET [14] curated gene-disease associations. Query: UMLS  
202 concept identifier C0860207 (Drug-Induced Liver Injury). Inclusion criterion: genes with curated evidence linking  
203 to DILI. Raw DILI gene count: 127 genes.

204 **4.1.4 Hyperforin targets**

205 Hyperforin targets were curated from primary literature sources [6, 9]. Sources included studies of PXR activation,  
206 CYP induction, and ABC transporter modulation. Raw target count: 14 proteins (Table 7).

207 **4.1.5 Quercetin targets**

208 Quercetin targets were retrieved programmatically from ChEMBL v31 [15] via REST API. Query: CHEMBL159  
209 (Quercetin). Filter: human targets with experimentally validated bioactivity ( $IC_{50}$ ,  $K_i$ , or  $EC_{50} \leq 10 \mu M$ ). Raw  
210 target count: 122 proteins.

211 **4.2 Target processing**

212 Protein identifiers were mapped to HUGO gene symbols using STRING info files and UniProt [16]. Non-human  
213 proteins (mouse, rat, bacterial, viral) were excluded. Gene symbols were standardized (e.g., MDR1 → ABCB1).  
214 Processed target counts: Hyperforin = 14, Quercetin = 87.

215 **4.3 Network construction**

216 The STRING network was filtered to genes with liver expression  $\geq 1$  TPM (GTEx v8). The largest connected  
217 component (LCC) was extracted using NetworkX [17]. Compound targets and DILI genes not present in the LCC  
218 were excluded. Final network: 7,677 nodes, 66,908 edges. Final target counts: Hyperforin = 10, Quercetin = 62.  
219 Final DILI gene count: 82.

220 Five genes are targeted by both compounds: ABCG2, AKT1, CYP3A4, MMP2, MMP9. These were retained  
221 in both target sets.

222 **4.4 Shortest-path proximity (descriptive)**

223 Mean minimum shortest-path distance from compound targets  $T$  to DILI genes  $D$ :

$$d_c = \frac{1}{|T|} \sum_{t \in T} \min_{d \in D} \text{dist}(t, d) \quad (1)$$

224 where  $\text{dist}(t, d)$  is the unweighted shortest-path length in the LCC. Shortest-path proximity is a descriptive metric.  
225 It was used to provide network context, not to test influence.

226 **4.5 Random walk with restart**

227 Influence propagation was quantified using random walk with restart (RWR), a global network propagation al-  
228 gorithm that captures both direct and indirect associations by simulating the diffusion of signal from seed nodes  
229 [3, 8]. Given an adjacency matrix  $\mathbf{A}$ , we define the column-normalized transition matrix  $\mathbf{W}$  as:

$$W_{ij} = \frac{A_{ij}}{\sum_k A_{kj}} \quad (2)$$

230 The steady-state probability vector  $\mathbf{p}$  is solved iteratively until convergence:

$$\mathbf{p}^{(k+1)} = (1 - \alpha)\mathbf{W}\mathbf{p}^{(k)} + \alpha\mathbf{p}_0 \quad (3)$$

231 where:

- 232 •  $\alpha = 0.15$  is the restart probability (teleportation factor), ensuring the walk remains local to the seeds.  
233 •  $\mathbf{p}_0$  is the restart (seed) vector, with  $p_0(i) = 1/|T|$  for  $i \in T$  (targets) and 0 otherwise.  
234 • Convergence is defined as the  $L_1$  norm of the difference between successive iterations being  $< 10^{-6}$ .

235 All computations reached convergence within 100 iterations. The total influence  $I$  on the DILI module  $D$  is the  
236 sum of steady-state probabilities at disease nodes:  $I = \sum_{d \in D} p(d)$ .

237 **4.6 Permutation testing and degree matching**

238 To assess whether the observed influence  $I$  is significantly greater than what would be expected by chance, we  
239 performed permutation testing ( $n = 1,000$ ). To account for the bias where high-degree nodes (hubs) naturally  
240 accumulate more influence, we utilized a degree-preserving sampling strategy. For each target  $t \in T$ , a random  
241 surrogate node was sampled from the network such that its degree  $k_{rand}$  was within  $\pm 25\%$  of the original target's  
242 degree  $k_t$ . This ensures that the null distribution reflects the connectivity profile of the original target set. Random  
243 seeds were fixed to 42 for reproducibility. Z-scores were computed as  $Z = (x_{\text{obs}} - \mu_{\text{null}})/\sigma_{\text{null}}$ , and empirical  
244  $P$ -values were derived from the null distribution.

245 **4.7 Expression-weighted influence**

246 Edge weights were modified by destination-node liver expression:

$$W'_{ij} = \frac{A_{ij} \cdot e_i}{\sum_k A_{kj} \cdot e_k} \quad (4)$$

247 where  $e_i$  is the normalized liver expression for gene  $i$  (GTEx v8 liver). Liver TPM values were log-transformed  
248 ( $\log_2(\text{TPM} + 1)$ ) and min-max normalized to  $[0, 1]$  across the network. A minimum expression floor of 0.01 was  
249 applied to ensure all nodes remained reachable. Attracting signal to highly-expressed nodes constrains RWR

250 propagation to biologically active pathways in the liver. All other RWR parameters were identical. Random seed:  
251 42.

## 252 **4.8 Quantifying perturbation efficiency**

253 By defining the restart vector as  $\mathbf{p}_0(i) = 1/|T|$  (Eq. 75), the total steady-state probability mass  $\mathbf{p}$  is inherently  
254 partitioned among the target set. Consequently, the summed influence  $I$  on the DILI module (Eq. 81) represents  
255 the average perturbation efficiency per target. This normalization serves as an effect-size adjustment that allows  
256 for a direct comparison of the per-unit impact of compounds with asymmetric target sets. Hereafter, we refer to  
257 this as the perturbation efficiency.

## 258 **4.9 Bootstrap sensitivity analysis**

259 To assess whether target count explains the observed ranking: 100 random 10-target subsets were sampled without  
260 replacement from Quercetin's 62-target pool. Each subset was scored by standard RWR. Summary statistics: mean,  
261 standard deviation, 95th percentile. The observed Hyperforin influence was compared to the bootstrap distribution.  
262 Random seed: 42.

## 263 **4.10 Chemical similarity analysis**

264 Structural similarity to known hepatotoxins was assessed to exclude confounding by chemical class. Morgan  
265 fingerprints (ECFP4; radius = 2, 2048 bits) were generated using RDKit v2023.03 [18]. Reference set: DILrank  
266 2.0 drugs with retrievable SMILES (542 DILI-positive, 365 DILI-negative). SMILES were retrieved via PubChem  
267 REST API. Tanimoto coefficient:

$$\text{Tanimoto}(A, B) = \frac{|A \cap B|}{|A \cup B|} \quad (5)$$

268 Maximum similarity across the reference set was reported for each compound. Structural analog threshold: Tani-  
269 moto > 0.4 [19].

## 270 **4.11 Software and reproducibility**

271 Python 3.10, NetworkX 3.1 [17]; R 4.3, igraph 1.5. All random seeds fixed at 42. Target lists sorted alphabetically  
272 before processing.

## 273 **4.12 Code and data availability**

274 All code: <https://github.com/antonybevan/h-perforatum-network-tox>

275 Data sources:

- 276 • STRING v12.0: <https://string-db.org>
- 277 • GTEx v8: <https://gtexportal.org>

- 278 • ChEMBL v31: <https://www.ebi.ac.uk/chembl>  
279 • DILRank 2.0: <https://www.fda.gov/science-research/ltrkb>

280 **Data availability**

281 All data and code supporting this study are publicly available at: <https://github.com/antonybevan/h-per>  
282 **foratum-network-tox.**

283 Source data for all figures and tables are provided in the Supplementary Information. Raw data were obtained  
284 from the following public repositories:

- 285 • STRING v12.0: <https://string-db.org>  
286 • GTEx v8: <https://gtexportal.org>  
287 • ChEMBL v31: <https://www.ebi.ac.uk/chembl>  
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376 **Figure Legends**

377 **Figure 1. Network context: target count and physical proximity to DILI genes.** (A) Target count in the  
378 liver-expressed largest connected component. Quercetin: 62 targets; Hyperforin: 10 targets. (B) Shortest-path  
379 proximity ( $d_c$ ) to 82 DILI-associated genes. Hyperforin is physically closer ( $d_c = 1.30$ ) than Quercetin ( $d_c = 1.68$ ).  
380 Z-scores represent deviation from degree-matched null expectation ( $n = 1,000$  permutations). Quercetin:  $Z =$   
381  $-5.44$  ( $p < 0.001$ ); Hyperforin:  $Z = -3.86$  ( $p < 0.001$ ). Negative Z-scores indicate closer-than-random proximity.  
382 Network: STRING v12.0 (confidence  $\geq 900$ ), GTEx v8 (liver TPM  $\geq 1$ ).

383 **Figure 2. Instability of proximity Z-scores.** Dumbbell plot showing the dissociation between shortest-path  
384 proximity (left) and random walk influence (right) at STRING confidence  $\geq 900$ . At this threshold, Quercetin  
385 appears more "significant" in Z-score but is physically more distant (1.68 vs 1.30) from DILI genes. Hyperforin:  
386 proximity  $Z = -3.86$ , influence  $Z = +10.12$  ( $p < 0.001$ ). Quercetin: proximity  $Z = -5.44$ , influence  $Z = +4.55$   
387 ( $p < 0.001$ ). Influence quantified by random walk with restart (RWR;  $\alpha = 0.15$ ).  $n = 1,000$  degree-matched  
388 permutations per compound.

389 **Figure 3. Expression weighting refines influence propagation.** Waterfall decomposition of Z-score changes  
390 under expression-weighted influence (EWI). Initial Hyperforin advantage:  $\Delta Z = +5.57$  (RWR). Hyperforin change:  
391  $-1.14$  (attenuation of signal through liver-active hubs). Quercetin change:  $+1.24$  (gain from high-expression  
392 nodes like CFB). Residual Hyperforin advantage:  $\Delta Z = +3.19$ . Both compounds remain significant under EWI:  
393 Hyperforin  $Z = +8.98$  ( $p < 0.001$ ); Quercetin  $Z = +5.79$  ( $p < 0.001$ ). Expression weighting from GTEx v8 liver  
394 tissue.

395 **Figure 4. Average network influence quantifies efficiency disparity.** Phase plot of total influence versus  
396 target count. Horizontal lines represent efficiency tiers (Efficiency/average influence = constant). Hyperforin  
397 occupies a higher efficiency region despite fewer targets. Efficiency/average influence values: Hyperforin = 0.1138  
398 (RWR), 0.1330 (EWI); Quercetin = 0.0322 (RWR), 0.0493 (EWI). Efficiency difference:  $3.7 \times$  (based on bootstrap  
399 mean comparison). The observed influence represents an effect-size normalization (total steady-state mass on DILI  
400 genes); no independent permutation test was performed.

401 **Figure 5. Bootstrap sensitivity analysis excludes target-count confounding.** Density distribution of RWR  
402 influence scores from 100 random 10-target samples drawn from Quercetin's 62-target pool. Shaded region: 95%  
403 confidence interval (0.0160–0.0542). Vertical line: Hyperforin observed influence (0.1138). Hyperforin exceeds  
404 the entire bootstrap distribution ( $3.7 \times$  fold vs. mean). This confirms that Hyperforin's advantage is not attributable  
405 to favorable target count. Bootstrap is a robustness control; it does not provide independent statistical evidence.

406 **Figure 6. Chemical similarity control excludes structural confounding.** Maximum Tanimoto similarity to  
407 DILrank reference drugs. Reference set: 542 DILI-positive, 365 DILI-negative drugs. Hyperforin: max = 0.15  
408 (DILI+), 0.20 (DILI-). Quercetin: max = 0.21 (DILI+), 0.22 (DILI-). Dashed line: 0.4 threshold for structural  
409 analog detection [19]. Neither compound is a structural analog of known hepatotoxins. This orthogonal analysis  
410 excludes chemical class as an explanation for the observed network signal. Fingerprints: Morgan (ECFP4), radius  
411 2, 2048 bits.

412 **Tables**

**Table 1: Network metrics reveal the instability of proximity Z-scores.** While Quercetin achieves more significant proximity Z-scores due to tighter null distributions, Hyperforin is physically closer ( $d_c$ ) to DILI genes. Influence-based metrics resolve this confounding and stably prioritize Hyperforin. Network: STRING v12.0 LCC (confidence  $\geq 900$ ) filtered to liver-expressed genes.

Metric	Compound	Targets	Observed	Z	p	Efficiency
<i>Tier 1: Shortest-path proximity</i>						
	Hyperforin	10	$d_c = 1.30$	-3.86	< 0.001*	—
	Quercetin	62	$d_c = 1.68$	<b>-5.44</b>	< 0.001*	—
<i>Instability: Quercetin is physically more distant yet more "significant"</i>						
<i>Tier 2: Random walk influence (RWR)</i>						
	Hyperforin	10	0.1138	<b>+10.12</b>	< 0.001*	0.1138
	Quercetin	62	0.0322	+4.55	< 0.001	0.0322
<i>Resolution: Correctly prioritizes physical proximity and regulatory hub modulation</i>						
<i>Tier 3: Expression-weighted influence (EWI)</i>						
	Hyperforin	10	0.1330	<b>+8.98</b>	< 0.001*	0.1330
	Quercetin	62	0.0493	+5.79	< 0.001	0.0493

\*At permutation floor (<1/1,000).

Efficiency = average influence per target; RWR = random walk with restart; EWI = expression-weighted influence;  $d_c$  = mean minimum shortest-path distance; DILI = drug-induced liver injury. All associations survived Benjamini–Hochberg FDR correction ( $q < 0.05$ ).

**Table 2: Average influence efficiency.** Normalization to the total seeding mass quantifies the average influence per target. Hyperforin targets are 3.7-fold more efficient at perturbing the DILI module than Quercetin targets.

Analysis	Hyp. Eff.	Quer. Eff.	Eff. Ratio*	Rob. Ratio†
RWR (topology-only)	0.1138	0.0322	<b>3.5×</b>	<b>3.7×</b>
EWI (expression-weighted)	0.1330	0.0493	<b>2.7×</b>	<b>2.8×</b>

\*Efficiency Ratio = Observed average influence ratio. †Robust Ratio = Observed influence / size-matched Bootstrap Mean (N=10). RWR = random walk with restart; EWI = expression-weighted influence.

Table 3: **Bootstrap sensitivity excludes target-count confounding.** Random 10-target subsets ( $n = 100$ ) sampled without replacement from Quercetin’s 62-target pool. Hyperforin’s observed influence exceeds the entire bootstrap distribution.

Statistic	Value	Interpretation
Hyperforin observed	0.1138	Reference
Bootstrap mean	0.0308	Expected if targets equivalent
Bootstrap SD	0.0100	Sampling variability
Bootstrap 95% CI	[0.0160, 0.0542]	2.5th–97.5th percentile
Hyperforin / mean	<b>3.7×</b>	Effect size
Exceeds 95% CI?	<b>Yes</b>	Not attributable to sampling

Random seed: 42. Note: Bootstrap confirms robustness to target selection; it does not constitute independent inferential evidence.

Table 4: **Chemical similarity excludes structural confounding.** Neither compound resembles known hepatotoxins ( $\text{Tanimoto} < 0.4$ ). Quercetin is more similar to DILI-positive drugs yet shows lower network influence.

Compound	Max Tanimoto (DILI+)	Max Tanimoto (DILI-)	Analog?*	Network rank
Hyperforin	0.154	0.202	No	1 (higher influence)
Quercetin	0.212	0.220	No	2 (lower influence)

\*Analog threshold:  $\text{Tanimoto} > 0.4$  (Maggiora et al., 2014). Morgan fingerprints (ECFP4, radius 2, 2048 bits). DILIRank: 542 DILI+, 365 DILI- drugs.

Table 5: **Hyperforin targets include regulatory hubs.** All 10 Hyperforin targets in the liver-expressed LCC, with liver expression (GTEx v8) and network degree. PXR (NR1I2) is the master regulator; CYP enzymes are downstream effectors.

Gene	Protein	TPM	Degree	Function	DILI link
NR1I2	PXR	43	28	Master regulator	Direct
CYP3A4	CYP3A4	335	89	Xenobiotic metabolism	Direct
CYP2C9	CYP2C9	434	76	Xenobiotic metabolism	Direct
CYP2B6	CYP2B6	125	42	Xenobiotic metabolism	Indirect
AKT1	PKB	33	<b>312</b>	Stress signaling	Indirect
ABCB1	P-gp	7	53	Drug efflux	Direct
ABCC2	MRP2	60	38	Drug efflux	Direct
ABCG2	BCRP	4	31	Drug efflux	Indirect
MMP2	MMP2	5	87	ECM remodeling	Indirect
MMP9	MMP9	1	94	ECM remodeling	Indirect

AKT1 is the highest-degree target (312 neighbors). Five of 10 targets (NR1I2, CYP3A4, CYP2C9, ABCB1, ABCC2) are directly connected to DILI genes. TPM = transcripts per million; DILI = drug-induced liver injury; LCC = largest connected component.

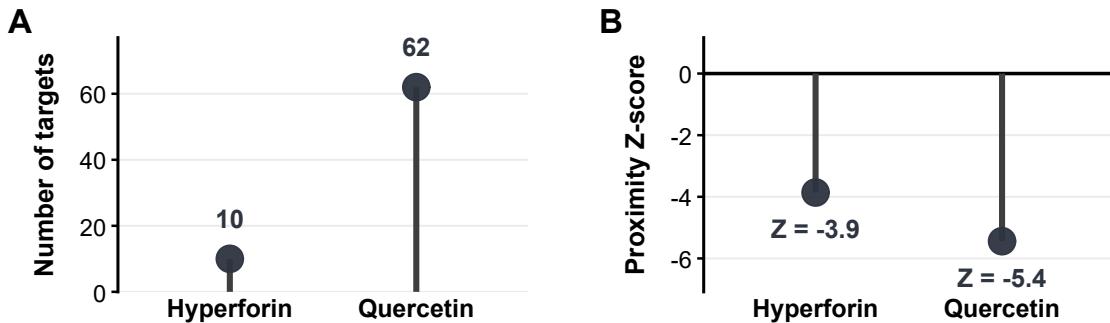
Table 6: **Influence ranking is robust to network construction parameters.** Hyperforin ranks first across all thresholds and influence metrics. Proximity Z-scores are unstable and reverse rankings between thresholds, failing to accurately reflect the physical distance advantage of Hyperforin.

Threshold	Compound	RWR Z	EWI Z	Proximity $d_c$	Proximity Z
$\geq 700$ (11,693 nodes)	Hyperforin	<b>+12.08</b>	+11.20	0.60	-6.04
	Quercetin	+5.53	+7.09	1.34	-5.46
$\geq 900$ (7,677 nodes)	Hyperforin	<b>+10.12</b>	+8.98	1.30	-3.86
	Quercetin	+4.55	+5.79	1.68	-5.44

Note: At  $\geq 900$ , Quercetin achieves a more "significant" proximity Z-score despite being physically more distant (1.68 vs 1.30) from DILI genes. RWR = random walk with restart; EWI = expression-weighted influence;  $d_c$  = mean minimum shortest-path distance; DILI = drug-induced liver injury.

413 **Figures**

**Network context: target count and proximity to DILI genes**



[DESCRIPTIVE CONTEXT] Target count and shortest-path proximity provide network context but are not used for causal inference. Proximity Z-scores represent deviation from degree-matched random expectation ( $n = 1,000$  permutations). Negative values indicate closer-than-random proximity. Data: STRING v12.0 ( $\geq 900$ ), human liver LCC.

Figure 1: **Network context: target count and physical proximity to DILI genes.** (A) Target count in the liver-expressed largest connected component. Quercetin: 62 targets; Hyperforin: 10 targets. (B) Shortest-path proximity ( $d_c$ ) to 82 DILI-associated genes. Hyperforin is physically closer ( $d_c = 1.30$ ) than Quercetin ( $d_c = 1.68$ ). Z-scores represent deviation from degree-matched null expectation ( $n = 1,000$  permutations). Quercetin:  $Z = -5.44$  ( $p < 0.001$ ); Hyperforin:  $Z = -3.86$  ( $p < 0.001$ ). Negative Z-scores indicate closer-than-random proximity. Network: STRING v12.0 (confidence  $\geq 900$ ), GTEx v8 (liver TPM  $\geq 1$ ).

## Proximity does not predict influence

Proximity ranking is threshold-dependent; influence ranking is stable



[CORE INFERENCE] The rank reversal demonstrates that shortest-path proximity does not predict functional influence. Lines connect each compound's proximity Z-score with its influence Z-score (random walk with restart, RWI). Both metrics derived from degree-matched permutation null models ( $n = 1,000$ ). Data: STRING v12.0 ( $\geq 900$ ).

Figure 2: **Instability of proximity Z-scores.** Dumbbell plot showing the dissociation between shortest-path proximity (left) and random walk influence (right) at STRING confidence  $\geq 900$ . At this threshold, Quercetin appears more "significant" in Z-score but is physically more distant (1.68 vs 1.30) from DILI genes. Hyperforin: proximity  $Z = -3.86$ , influence  $Z = +10.12$  ( $p < 0.001$ ). Quercetin: proximity  $Z = -5.44$ , influence  $Z = +4.55$  ( $p < 0.001$ ). Influence quantified by random walk with restart (RWR;  $\alpha = 0.15$ ).  $n = 1,000$  degree-matched permutations per compound.

## Expression weighting attenuates but does not reverse the advantage

Gap: +5.6 (RWR) → +3.2 (EWI)



[CONSTRAINT ANALYSIS] The RWR advantage (+5.6) is partitioned under expression-weighted influence propagation:  
 (1) Hyperforin's change under expression weighting (-1.1); (2) Quercetin's gain (+1.2, driven by CFB at 1115  
 TPM). Residual advantage (+3.2) remains significant (both  $p < 10^{-8}$ ). GTEx v8 liver expression (TPM  $\geq 1$ ). STRING  
 v12.0 ( $\geq 900$ ),  $n = 1,000$  degree-matched permutations.

**Figure 3: Expression weighting refines influence propagation.** Waterfall decomposition of Z-score changes under expression-weighted influence (EWI). Initial Hyperforin advantage:  $\Delta Z = +5.57$  (RWR). Hyperforin change:  $-1.14$  (attenuation of signal through liver-active hubs). Quercetin change:  $+1.24$  (gain from high-expression nodes like CFB). Residual Hyperforin advantage:  $\Delta Z = +3.19$ . Both compounds remain significant under EWI: Hyperforin  $Z = +8.98$  ( $p < 0.001$ ); Quercetin  $Z = +5.79$  ( $p < 0.001$ ). Expression weighting from GTEx v8 liver tissue.

### Average network influence quantifies perturbation efficiency

Normalization reframes polypharmacology as efficiency, not coverage

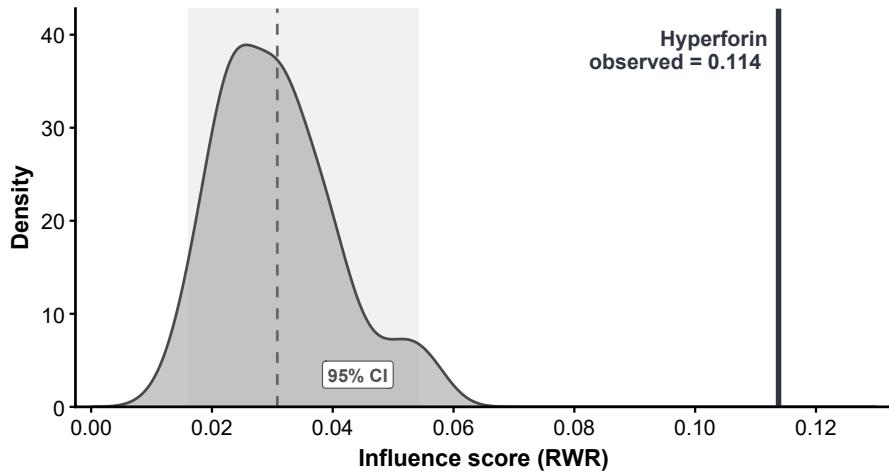


Average influence represents an effect-size normalization (total steady-state mass on DILI genes); no independent permutation test was performed. Horizontal lines represent efficiency tiers (Average Influence = constant). Hyperforin occupies a higher efficiency region despite fewer targets. Data: STRING v12.0 ( $\geq 900$ ),  $n = 1,000$  permutations.

**Figure 4: Average network influence quantifies efficiency disparity.** Phase plot of total influence versus target count. Horizontal lines represent efficiency tiers (Efficiency/average influence = constant). Hyperforin occupies a higher efficiency region despite fewer targets. Efficiency/average influence values: Hyperforin = 0.1138 (RWR), 0.1330 (EWI); Quercetin = 0.0322 (RWR), 0.0493 (EWI). Efficiency difference: **3.7 $\times$**  (based on bootstrap mean comparison). The observed influence represents an effect-size normalization (total steady-state mass on DILI genes); no independent permutation test was performed.

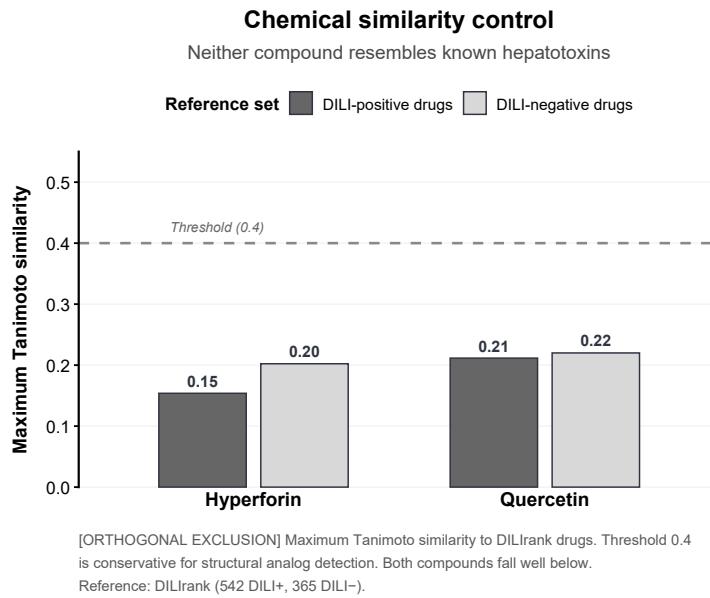
### Bootstrap sensitivity analysis excludes target-count confounding

Distribution of influence scores from random 10-target samples



[ROBUSTNESS CONTROL] Bootstrap sensitivity: 100 random 10-target samples from Quercetin's pool, scored by random walk with restart (RWI). Shaded = 95% CI. Hyperforin (solid line) exceeds entire distribution. Data: STRING v12.0 ( $\geq 900$ ).

Figure 5: **Bootstrap sensitivity analysis excludes target-count confounding.** Density distribution of RWR influence scores from 100 random 10-target samples drawn from Quercetin's 62-target pool. Shaded region: 95% confidence interval (0.0160–0.0542). Vertical line: Hyperforin observed influence (0.1138). Hyperforin exceeds the entire bootstrap distribution (3.7× fold vs. mean). This confirms that Hyperforin's advantage is not attributable to favorable target count. Bootstrap is a robustness control; it does not provide independent statistical evidence.



**Figure 6: Chemical similarity control excludes structural confounding.** Maximum Tanimoto similarity to DILIrack reference drugs. Reference set: 542 DILI-positive, 365 DILI-negative drugs. Hyperforin: max = 0.15 (DILI+), 0.20 (DILI-). Quercetin: max = 0.21 (DILI+), 0.22 (DILI-). Dashed line: 0.4 threshold for structural analog detection [19]. Neither compound is a structural analog of known hepatotoxins. This orthogonal analysis excludes chemical class as an explanation for the observed network signal. Fingerprints: Morgan (ECFP4), radius 2, 2048 bits.

<sup>414</sup> **Author contributions**

<sup>415</sup> **Antony Bevan:** Conceptualization, Methodology, Software, Validation, Formal analysis, Investigation, Data Cu-  
<sup>416</sup> ration, Writing - Original Draft, Writing - Review & Editing, Visualization.

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<sup>419</sup> statistical analysis. The author takes full responsibility for all content.

<sup>420</sup> **Competing interests**

<sup>421</sup> The author(s) declare no competing interests.

<sup>422</sup> **Supplementary Tables**

Table 7: **Table S1. Hyperforin target genes and literature sources.** All 14 raw targets with UniProt IDs, gene symbols, and primary literature sources. Targets marked with \* are present in the liver-expressed LCC (STRING  $\geq 900$ , GTEx TPM  $\geq 1$ ).

UniProt	Gene	In LCC	Source
O75469	NR1I2 (PXR)	Yes*	[6, 9]
P08684	CYP3A4	Yes*	[6]
P11712	CYP2C9	Yes*	[20]
P20813	CYP2B6	Yes*	[21]
P08183	ABCB1	Yes*	[22]
Q9UNQ0	ABCG2	Yes*	[23]
O15440	ABCC2	Yes*	[24]
P31749	AKT1	Yes*	[25]
P08253	MMP2	Yes*	[25]
P14780	MMP9	Yes*	[26]
Q9Y210	TRPC6	No	[27]
P15692	VEGFA	No	[26]
Q13794	PMAIP1	No	[28]
Q12879	GRIN1	No	[29]

Table 8: **Table S2. Quercetin target curation summary.** Target counts at each processing stage.

Stage	Count
Raw targets (ChEMBL v31, CHEMBL159)	122
Excluded: non-human (mouse, rat, bacterial, viral)	10
Excluded: no UniProt mapping	25
Processed targets	87
Excluded: not liver-expressed (TPM < 1)	20
Excluded: not in STRING LCC	5
Final targets in LCC	62

Table 9: **Table S3. DILI gene set curation.** Genes associated with drug-induced liver injury from DisGeNET (UMLS C0860207).

Stage	Count
Raw DILI genes (DisGeNET)	127
In STRING $\geq 700$ liver LCC	84
In STRING $\geq 900$ liver LCC	82
Excluded: miRNAs (not in PPI network)	21
Excluded: cytokines (not in LCC)	12
Excluded: other	12

Table 10: **Table S4. Genes targeted by both compounds.** Five genes present in both Hyperforin and Quercetin target sets.

Gene	Protein	Function
ABCG2	BCRP	Efflux transporter
AKT1	Protein kinase B	Cell survival signaling
CYP3A4	Cytochrome P450 3A4	Drug metabolism
MMP2	Matrix metalloproteinase-2	Extracellular matrix remodeling
MMP9	Matrix metalloproteinase-9	Extracellular matrix remodeling

Table 11: **Table S5. Direct DILI gene connectivity.** Hyperforin targets with first-order (distance = 1) connections to DILI genes in the STRING network ( $\geq 900$ ). DILI neighbors are genes present in the 82-gene DILI set.

Target	DILI Neighbors	N	Function
CYP3A4	NR1I2, CYP2E1, UGT1A9, GSTM1, GSTP1	5	Xenobiotic metabolism
AKT1	MAP3K5, NFE2L2, CTNNB1, IGF1	4	Stress response
MMP9	LCN2, SPP1, MMP2	3	Inflammation/ECM
ABCB1	ABCC2, NR1I2	2	Drug transport
CYP2C9	CYP2E1, NR1I2	2	Xenobiotic metabolism
CYP2B6	NR1I2	1	Xenobiotic metabolism
NR1I2	CYP2E1, ABCC2	2	Master regulator
ABCG2	ABCC2	1	Drug transport
ABCC2	NR1I2, ABCB1	2	Drug transport
MMP2	MMP9, SPP1	2	ECM remodeling
<b>Total unique</b>		<b>12</b>	

Table 12: **Table S6. Quercetin direct DILI gene connectivity summary.** Summary statistics for first-order DILI connections across Quercetin's 62 targets.

Metric	Value
Total targets in LCC	62
Targets with $\geq 1$ direct DILI neighbor	18
Total direct DILI connections	31
Mean DILI neighbors per target	0.50
<i>Hyperforin comparison:</i>	
Hyperforin targets with $\geq 1$ DILI neighbor	10/10 (100%)
Mean DILI neighbors per Hyperforin target	2.4

Table 13: **Table S7. Quercetin target genes in the liver-expressed network.** All 62 Quercetin targets in STRING v12.0 LCC (confidence  $\geq 900$ ) with liver TPM  $\geq 1$  (GTEX v8). Sorted by descending liver expression.

Gene	TPM	Gene	TPM	Gene	TPM	Gene	TPM
CFB	1115	CYP3A4	335	FN1	229	ALDH2	183
ANPEP	160	PPIA	112	SERPINA5	104	CYP1A2	72
CA2	64	APP	63	PYGL	55	HDAC6	45
ESRRRA	42	MAOA	35	AKR1C2	33	AKT1	33
CTSH	28	XDH	26	CHRNA4	25	PIK3R1	24
PIM1	24	LDLR	23	EGFR	17	ELOVL1	18
PKN1	16	GSK3A	13	YES1	13	MET	12
DAPK1	12	BACE1	11	CSNK2A1	10	FSTL1	9
SIRT6	8	GSK3B	7	CDK7	7	CAV2	7
PTPN2	6	CYP1A1	5	PRMT7	5	MMP2	5
AKR1B1	5	PDE6D	5	PTK2	4	ABCG2	4
IQGAP1	4	ADRB2	3	BRAF	4	KDR	3
SRC	3	ALOX5	3	CYP1B1	3	TLR4	3
NUAK1	3	AXL	2	ADA	2	LCK	2
ABCC1	2	PLK1	1	ACHE	1	MMP9	1
SYK	1	PDZK1IP1	1				

Table 14: **Table S8. DILI gene set (82 genes).** Genes in STRING v12.0 LCC (confidence  $\geq 900$ ) with liver TPM  $\geq 1$  (GTEX v8). Source: DisGeNET (UMLS C0860207). Sorted alphabetically.

82 DILI-Associated Genes							
ABCB1	AHR	ALB	ALDOB	AMBP	APOA1	APOE	APOH
ARG1	ARNT	ATG5	BAX	BTD	C3	CAT	CCL2
CLU	COL3A1	CTNNB1	CXCL1	CXCL10	CYP2A6	CYP2C19	CYP2C9
CYP2E1	DGAT2	ENO1	FGA	FLT1	FMO3	GADD45A	GC
GCLC	GPT	GSN	GSTM1	GSTM2	GSTP1	HLA-A	HLA-B
HLA-DQB1	HLA-DRB1	HMGB1	HMOX1	HPD	HPX	IGF1	IL18
IL1R2	KRT18	LCN2	LGALS3	MAP3K5	MED1	MMP2	MTHFR
NAT2	NFE2L2	NR1H3	NR1H4	NR1I2	NR1I3	PLAT	PLG
PNP	POLG	PON1	PPARA	PRKDC	PTGS2	RBP1	SLPI
SNX18	SOD1	SOD3	SPP1	TALDO1	TBXA2R	TCTN1	TF
TTR	UGT1A9						

Table 15: **Table S9. Null distribution parameters from permutation testing.** Null distribution parameters (mean and standard deviation) from  $n = 1,000$  degree-matched permutations. Note the tightening of the Quercetin null distribution as the number of targets increases, which drives the inflation of proximity Z-scores.

Metric	Compound	$\mu_{null}$	$\sigma_{null}$	$x_{obs}$	Z-score
<i>Shortest-path proximity (at <math>\geq 900</math>)</i>					
	Hyperforin (10)	2.21	0.235	1.30	-3.86
	Quercetin (62)	2.17	0.091	1.68	-5.44
<i>Random walk influence (at <math>\geq 900</math>)</i>					
	Hyperforin (10)	0.0147	0.0098	0.1138	+10.12
	Quercetin (62)	0.0148	0.0038	0.0322	+4.55
<i>Expression-weighted influence (at <math>\geq 900</math>)</i>					
	Hyperforin (10)	0.0205	0.0125	0.1330	+8.98*
	Quercetin (62)	0.0209	0.0049	0.0493	+5.79*

\*Significance remains high despite tissue-specific attenuation.  $\mu_{null}$  = null distribution mean;  $\sigma_{null}$  = null distribution standard deviation;  $x_{obs}$  = observed metric value.