Perturbation efficiency resolves target-count bias in network proximity metrics: A controlled audit

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Abstract

Network-based metrics are essential for identifying drug–disease associations, assuming that proximity within protein–protein interaction networks reflects functional relevance. However, proximity Z-scores are fundamentally sensitive to target set size. This introduces a systematic bias where compounds with broad polypharmacology appear statistically significant due to null distribution tightening (the Law of Large Numbers) rather than physical reachability. We systematically audit this bias using the human liver interactome and constituents from *Hypericum perforatum*. We demonstrate that proximity Z-scores yield unstable rankings that reverse across network construction parameters. While compounds with many targets may achieve higher significance, they can remain physically more distant from disease modules than high-leverage modulators. We resolve this by utilizing random walk–based influence propagation and introducing *perturbation efficiency* to ensure unbiased comparisons. Our results show that influence-based rankings are stable and correctly identify high-leverage modulators that proximity metrics miss. This study provides a methodological template for identifying and correcting statistical artifacts in network medicine, enabling reliable assessments in complex biological systems.

Keywords: network medicine, proximity metrics, metric robustness, drug–induced liver injury, Z-score bias, perturbation efficiency.

1 Introduction

Network-based prioritization is a cornerstone of modern systems biology and drug discovery, assuming that the topological proximity between compound targets and disease genes within a protein–protein in-

teraction (PPI) network reflects functional relevance [1–4]. Because raw network distances are sensitive to local topology and degree distribution, they are typically reported as Z-scores relative to degree-matched null models. While these Z-scores successfully quantify statistical significance in most applications [3], we demonstrate they can be confounded by large asymmetries in target set size. As the number of seed nodes increases, the variance of the null distribution decreases (the Law of Large Numbers), leading to deterministic significance inflation for compounds with broad polypharmacology. Identifying whether such results reflect true biological influence, or whether they represent systematic artifacts of distance-based inference, is essential for the reliability of network medicine.

Using the human liver interactome as a model system, we investigate this confounding effect through a controlled comparison of two constituents from *Hypericum perforatum* (St. John's Wort). These constituents—Hyperforin and Quercetin—exhibit highly asymmetric target set sizes: Hyperforin possesses 10 validated targets, while Quercetin has over 60 [5–7]. This system serves as a controlled model because it pairs a known biological ground truth (Hyperforin-mediated PXR activation) with extreme topological asymmetry, providing a sharp stress test for network metrics. Conventional proximity Z-scores predict greater disease-associated significance for the broad-spectrum modulator, even when it is physically more distant from the disease module than the high-leverage modulator. This reversal indicates that proximity-based prioritization is unstable across network construction parameters and susceptible to sample-size artifacts.

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

2 Results

2.1 Proximity Z-scores are confounded by target set size

We first established network context by quantifying target count and shortest-path proximity to 82 DILI-associated genes (Figure 1). Quercetin engages 62 targets in the liver-expressed largest connected

component; Hyperforin engages 10. At STRING confidence ≥900, Hyperforin targets are physically closer to DILI genes ($d_c = 1.30$) than Quercetin targets ($d_c = 1.68$; Table 1). However, the proximity Z-scores yield the opposite ranking: Quercetin achieves $Z = -5.44$ ($p < 0.001$), while Hyperforin achieves $Z = -3.86$ ($p < 0.001$). All reported associations survived Benjamini–Hochberg FDR correction ($q < 0.05$).

This statistical artifact suggests that Quercetin poses greater risk, whereas the physical topology favors Hyperforin. This effect occurs independently of network construction parameters and represents a fundamental statistical property of averaged distributions: larger samples inherently produce more precise (narrower) null distributions, artificially inflating Z-score magnitude regardless of actual topological proximity. This is a manifestation of the Law of Large Numbers (LLN).

2.2 Influence-based rankings are stable and resolve the confound

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

2.3 Expression weighting refines the signal

To assess whether the RWR signal persists under tissue-specific constraint, we applied expression-weighted influence propagation (EWI), weighting transitions by destination-node liver expression (Figure 3).

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

2.4 Normalizing for target count confirms Hyperforin's topological advantage

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

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

*RWR Ratio: observed ratio (robust ratio in parentheses). †EWI Ratio: observed ratio (robust ratio in parentheses).

Each Hyperforin target contributes 3.7× more DILI-directed influence than each Quercetin target (robust ratio). This disparity indicates that Hyperforin's target positions are substantially higher leverage than those of Quercetin, achieving greater perturbation efficiency despite a 6-fold smaller target set. The efficiency ratio remains stable within a narrow range: 3.7× at STRING ≥700, 3.5× at ≥900 (a 5% variation despite 33% reduction in network density). In contrast, proximity Z-score rankings undergo complete reversal between these thresholds. This demonstrates that perturbation efficiency is a parameter-invariant comparative metric where distance-based Z-scores are not.

2.5 Bootstrap resampling excludes target-selection bias

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

Hyperforin's observed influence (0.1138) exceeds the entire bootstrap distribution from Quercetin (mean = 0.0308, 95% CI = [0.0160, 0.0542]; Supplementary Table S10). The fold difference between Hyperforin and the bootstrap mean is 3.7×. This confirms that Hyperforin's advantage is not an artifact of target count; even when sampling equalized subsets from Quercetin's pool, no configuration matches Hyperforin's influence.

2.6 Ranking stability across network thresholds

The influence ranking is stable across network confidence thresholds (Supplementary Table S13). Hyperforin ranks first in all RWR and EWI configurations at both ≥700 and ≥900 thresholds. Notably, the proximity ranking reverses between thresholds: at ≥700, Hyperforin is physically closer ($d_c = 0.60$ vs 1.34) and more "significant" ($Z = -6.04$ vs -5.46). At ≥900, Quercetin appears more "significant" ($Z = -5.44$ vs -3.86) despite being

physically more distant (1.68 vs 1.30). This instability in proximity Z-scores—while influence rankings remain stable—demonstrates that influence-based metrics are more robust to network construction parameters.

2.7 Chemical similarity excludes structural confounding

To exclude the possibility that Hyperforin's network signal reflects structural similarity to known hepatotoxins, we performed chemical similarity analysis against the DILIrank reference set (Figure 6). Morgan fingerprints (ECFP4) revealed that neither compound exceeds the 0.4 Tanimoto threshold for structural analog detection. Notably, Quercetin exhibits higher structural similarity to DILI reference drugs yet lower network influence, reinforcing that the observed asymmetry is driven by network topology rather than chemical features.

3 Discussion

3.1 Ranking stability and the Z-score confound

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

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

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

3.2 Relationship to prior work

Our findings do not contradict the foundational work of Guney et al. (2016), but rather identify a specific failure mode that their study design did not stress-test. Guney et al. evaluated network proximity as a classifier for drug-disease associations across 238 drugs with a mean of 3.5 targets per drug—a relatively homogeneous dataset. They reported that proximity is "not biased with respect to the number of targets a drug has" and found that the

closest-distance measure (d_c) outperformed a diffusion kernel measure (d_k) for binary classification [3].

Our study addresses a fundamentally different question: *comparative ranking* of two compounds with highly asymmetric target counts (10 vs. 62). In this regime, the variance-shrinkage artifact becomes a first-order problem. Guney's kernel benchmark (d_k) is related to but distinct from random walk with restart; d_k sums contributions from all weighted paths, whereas RWR iteratively propagates probability mass with a restart factor that anchors the walk to seed nodes. More critically, neither d_c nor d_k provides a principled normalization for target set size.

A primary methodological resolution proposed in this study is the use of *perturbation efficiency*: the average influence exerted per target. This normalization resolves the target-count paradox regardless of whether the underlying propagation method is shortest-path, kernel, or random walk. By framing polypharmacology as an efficiency problem rather than a coverage problem, we provide a bias-corrected comparative framework that survives robustness checks where raw Z-scores fail. While Guney et al. found that a diffusion kernel underperformed closest distance for binary classification of known drug-disease pairs, our task differs fundamentally: we address comparative ranking under extreme target-count asymmetry (10 vs. 62 targets). RWR's restart mechanism enforces locality absent in pure diffusion kernels, and our empirical results demonstrate stable rankings that align with biological ground truth—a criterion not assessed in Guney's benchmark.

3.3 Expression weighting as a biological constraint

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

3.4 Perturbation efficiency vs. topological coverage

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

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

3.5 Mechanistic context: The PXR axis

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

3.6 Limitations

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

Second, while we demonstrate that proximity Z-scores are confounded by target set size, influence-based Z-scores are not entirely immune to this effect. As the number of seed nodes increases, the variance of the null distribution for influence sums also decreases, though less severely than for distance-based metrics. Critically, our core claims do not rest on absolute Z-score comparisons. We demonstrate that influence-based *rankings* are stable across network thresholds, while proximity rankings are not. We further resolve the size-dependence by introducing perturbation efficiency (influence per target), which explicitly normalizes for target count and provides a bias-corrected comparative metric.

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

3.7 Conclusions

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

More broadly, this work provides a methodological template for identifying and resolving metric artifacts

in network toxicology. By integrating signed edge weights and transcriptometric data, future iterations of this framework could investigate phenotype-specific associations, linking topological influence on specific biological sub-modules to discrete clinical outcomes.

4 Methods

4.1 Data sources

4.1.1 Protein–protein interaction network

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

4.1.2 Liver expression data

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

4.1.3 Drug-induced liver injury gene set

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

4.1.4 Hyperforin targets

Hyperforin targets were curated from primary literature sources [6, 9]. Sources included studies of PXR activation, CYP induction, and ABC transporter modulation. Raw target count: 14 proteins (Supplementary Table S1).

4.1.5 Quercetin targets

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

4.2 Target processing

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

4.3 Network construction

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

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

4.4 Shortest-path proximity (descriptive)

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

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

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

4.5 Random walk with restart

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

$$W_{ij} = \frac{A_{ij}}{\sum_k A_{kj}} \tag{2}$$

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

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

where:

- $\alpha = 0.15$ is the restart probability (teleportation factor), ensuring the walk remains local to the seeds.
- $\mathbf{p}_0$ is the restart (seed) vector, with $p_0(i) = 1/|T|$ for $i \in T$ (targets) and 0 otherwise.

- Convergence is defined as the L_1 norm of the difference between successive iterations being $< 10^{-6}$.

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

4.6 Permutation testing and degree matching

To assess whether the observed influence I is significantly greater than what would be expected by chance, we performed permutation testing ($n = 1{,}000$). To account for the bias where high-degree nodes (hubs) naturally accumulate more influence, we utilized a degree-preserving sampling strategy. For each target $t \in T$, a random surrogate node was sampled from the network such that its degree k_{rand} was within $\pm 25\%$ of the original target's degree k_t. This ensures that the null distribution reflects the connectivity profile of the original target set. Random seeds were fixed to 42 for reproducibility. Z-scores were computed as $Z = (x_{\text{obs}} - \mu_{\text{null}})/\sigma_{\text{null}}$, and empirical P-values were derived from the null distribution. All statistical tests used a significance threshold of $\alpha = 0.05$ (two-tailed).

4.7 Expression-weighted influence

Edge weights were modified by destination-node liver expression:

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

where e_i is the normalized liver expression for gene i (GTEx v8 liver). Liver TPM values were log-transformed ($\log_2(\text{TPM} + 1)$) and min-max normalized to $[0, 1]$ across the network. A minimum expression floor of 0.01 was applied to ensure all nodes remained reachable. Attracting signal to highly-expressed nodes constrains RWR propagation to biologically active pathways in the liver. All other RWR parameters were identical. Random seed: 42.

4.8 Quantifying perturbation efficiency

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

4.9 Bootstrap sensitivity analysis

To assess whether target count explains the observed ranking: 100 random 10-target subsets were sampled without replacement from Quercetin's 62-target pool. Each subset was scored by standard RWR. Summary statistics: mean,

standard deviation, 95th percentile. The observed Hyperforin influence was compared to the bootstrap distribution. Random seed: 42.

4.10 Chemical similarity analysis

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

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

Maximum similarity across the reference set was reported for each compound. Structural analog threshold: Tanimoto > 0.4 [19].

4.11 Software and reproducibility

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

Use of AI tools

AI-assisted tools were used to assist with code development and statistical analysis. The author takes full responsibility for all content and conclusions presented in this study.

Data Availability

All data and code supporting this study are publicly available in the following repository: `https://github.com/antonybevan/h-perforatum-network-tox`

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

- STRING v12.0: `https://string-db.org`
- GTEx v8: `https://gtexportal.org`
- ChEMBL v31: `https://www.ebi.ac.uk/chembl`
- DisGeNET: `https://www.disgenet.org`
- DILIrank 2.0: `https://www.fda.gov/science-research/ltkb`

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Author contributions

Antony Bevan: Conceptualization, Methodology, Software, Validation, Formal analysis, Investigation, Data Curation, Writing - Original Draft, Writing - Review & Editing, Visualization.

Competing interests

The author declares no competing interests.

Additional information

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Figure Legends

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).

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.

Figure 3. Expression weighting refines influence propagation. Waterfall decomposition of Z-score changes under expression-weighted influence (EWI). Initial Hyperforin advantage: ΔZ = +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: ΔZ = +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.

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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$\times$ 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 DILIrank 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.

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*-score	*P*-value	Efficiency
Tier 1: Shortest-path proximity						
	Hyperforin	10	$d_c = 1.30$	-3.86	$< 0.001^*$	—
	Quercetin	62	$d_c = 1.68$	$\mathbf{-5.44}$	$< 0.001^*$	—
	Instability: Quercetin is physically more distant yet more "significant"					
Tier 2: Random walk influence (RWR)						
	Hyperforin	10	0.1138	**+10.12**	$< 0.001^*$	0.1138
	Quercetin	62	0.0322	+4.55	< 0.001	0.0322
	Resolution: Correctly prioritizes physical proximity and regulatory hub modulation					
Tier 3: Expression-weighted influence (EWI)						
	Hyperforin	10	0.1330	**+8.98**	$< 0.001^*$	0.1330
	Quercetin	62	0.0493	+5.79	< 0.001	0.0493

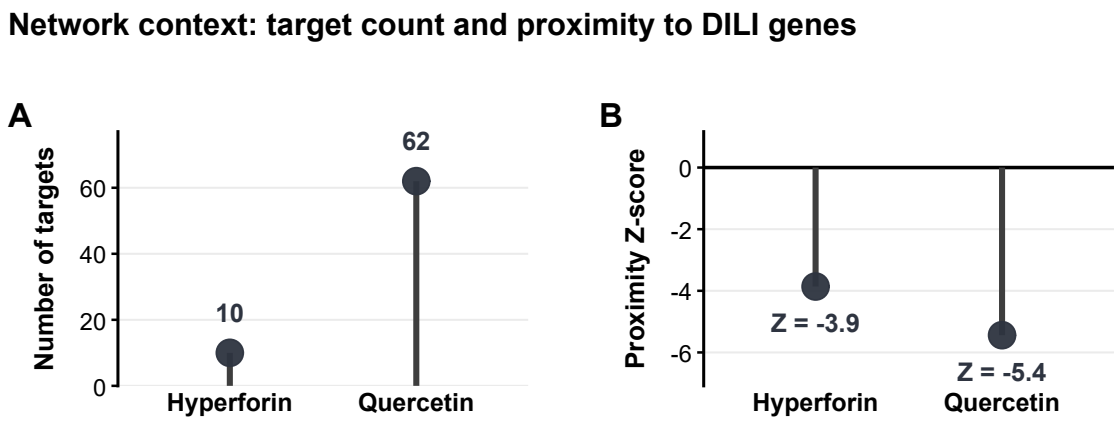
*At permutation floor ($<$1/1,000).
Efficiency = average influence per target; RWR = random walk with restart; EWI = expression-weighted influence; d_c = mean minimum shortest-path distance; DILI = drug-induced liver injury. All associations survived Benjamini–Hochberg FDR correction ($q < 0.05$).

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	**3.5**$\times$	**3.7**$\times$
EWI (expression-weighted)	0.1330	0.0493	**2.7**$\times$	**2.8**$\times$

*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.

Figures



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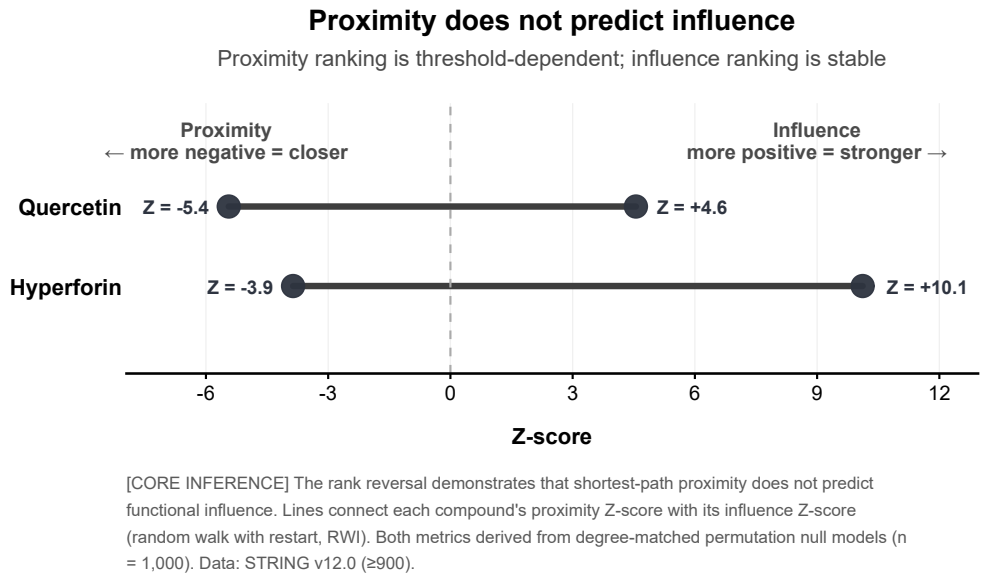

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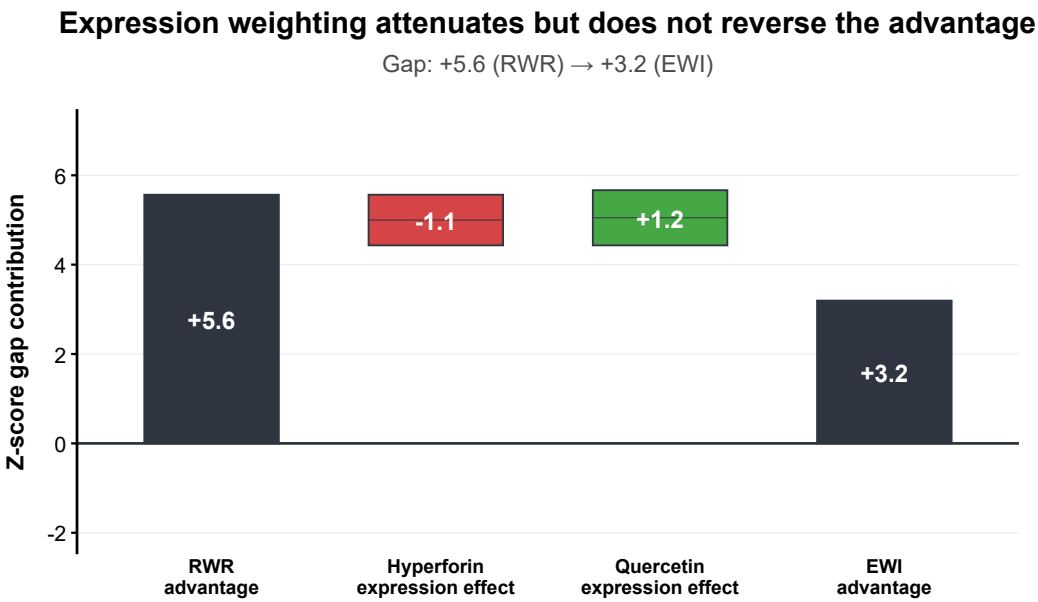

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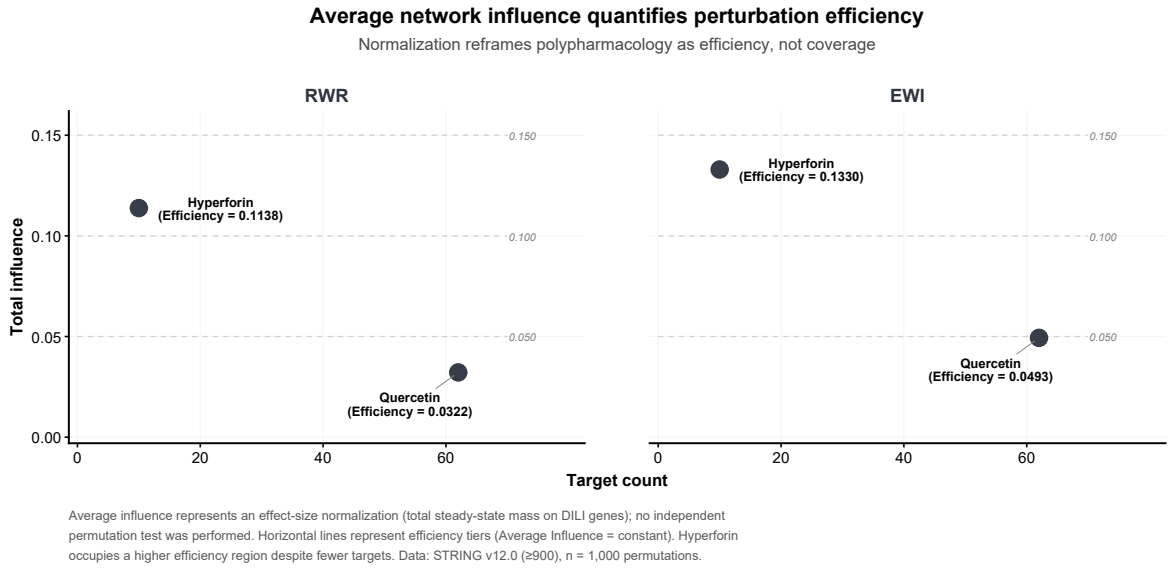

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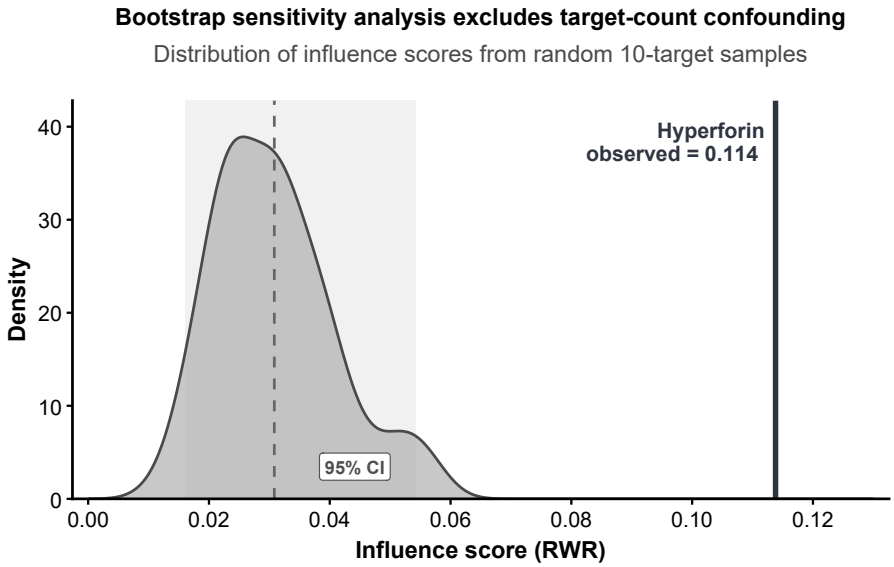

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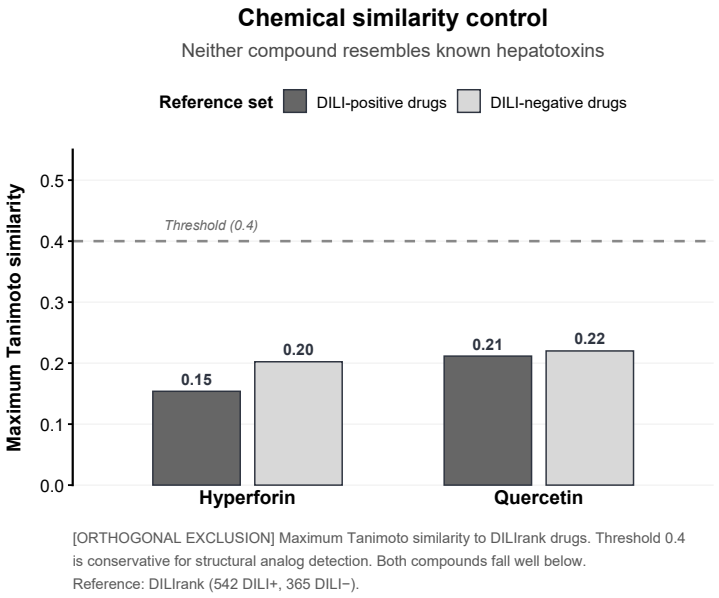

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Supplementary Information

Perturbation efficiency resolves target-count bias in network proximity metrics: A controlled audit

Antony Bevan

Supplementary Tables

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 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 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 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 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
Total unique		**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 ≥1 direct DILI neighbor	18
Total direct DILI connections	31
Mean DILI neighbors per target	0.50
Hyperforin comparison:	
Hyperforin targets with ≥1 DILI neighbor	10/10 (100%)
Mean DILI neighbors per Hyperforin target	2.4

Table S7: **Quercetin target genes in the liver-expressed network.** All 62 Quercetin targets in STRING v12.0 LCC (confidence ≥900) with liver TPM ≥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
ESRRA	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 S8: **DILI gene set (82 genes).** Genes in STRING v12.0 LCC (confidence ≥900) with liver TPM ≥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 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	μ_{null}	σ_{null}	x_{obs}	Z-score
Shortest-path proximity (at ≥900)					
	Hyperforin (10)	2.21	0.235	1.30	−3.86
	Quercetin (62)	2.17	0.091	1.68	−5.44
Random walk influence (at ≥900)					
	Hyperforin (10)	0.0147	0.0098	0.1138	+10.12
	Quercetin (62)	0.0148	0.0038	0.0322	+4.55
Expression-weighted influence (at ≥900)					
	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. μ_{null} = null distribution mean; σ_{null} = null distribution standard deviation; x_{obs} = observed metric value.

Table S10: **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	**3.7**×	Effect size
Exceeds 95% CI?	**Yes**	Not attributable to sampling

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

Table S11: **Chemical similarity excludes structural confounding.** Neither compound resembles known hepatotoxins (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: Tanimoto > 0.4 (Maggiora et al., 2014). Morgan fingerprints (ECFP4, radius 2, 2048 bits). DILIrank: 542 DILI+, 365 DILI− drugs.

Table S12: **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	**312**	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 S13: **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	Hyperforin	**+12.08**	+11.20	0.60	−6.04
(11,693 nodes)	Quercetin	+5.53	+7.09	1.34	−5.46
$\geq$900	Hyperforin	**+10.12**	+8.98	1.30	−3.86
(7,677 nodes)	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.