

Convert a reconstruction into a flux balance analysis model

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Reviewers:

INTRODUCTION

Even with quality control during the reconstruction process, it is not appropriate to assume that any reconstruction can be converted directly into a model and used to make predictions. A model must satisfy certain assumptions before it can be used to make reliable predictions. Depending on the type of model model, these assumptions will be different. Each assumption should be chemically or biologically motivated and expressed in an unambiguous manner and preferably both intuitively and mathematically. Flux balance analysis is a mathematical method widely used for studying genome-scale biochemical network. Here one aims to predict steady-state reaction fluxes, where there is a balance between production and consumption of each molecular species that is not exchanged across the specified boundary of a system. In this situation, one might obtain erroneous predictions if the system boundary is incorrectly specified. If a reconstruction contains one or more supposedly mass balanced reactions, but which are actually not mass balanced, such reactions in a model can lead to inadvertent leakage of a metabolite from the model, in violation of mass balance. Similarly, when generating a model for flux balance analysis, it is important to ensure that the network is flux consistent, that is, each reaction can carry a non-zero steady state flux.

Given a reconstruction with \hat{m} reactants involved in \hat{n} reactions, this tutorial demonstrates a method to identify and extract the largest subset of the reconstruction whose internal reactions are both stoichiometrically and flux consistent and whose external reactions are flux consistent. This model is then mathematically consistent with the basic requirements for generation of predictions using flux balance analysis. The identification of the component of the reconstruction that does not satisfy the aforementioned modelling conditions is also useful for targeting reconstruction effort towards resolving stoichiometric inconsistency or resolving flux inconsistency. The example used in this tutorial illustrates the process of extracting a model consistent with flux balance analysis, from a ReconX reconstruction.

PROCEDURE

Select reconstruction to convert into a model and enter parameters

Load the ReconX reconstruction, and save the original reconstruction in the workspace, unless it is already loaded into the workspace.

```
clear model
if ~exist('modelOrig','var')
    %select your own model, or use Recon2.0model instead
    if 1
        filename='Recon3D_301.mat'
        load(filename);
        model=Recon3D;
```

```

else
    filename='Recon2.0model.mat';
    if exist('Recon2.0model.mat','file')==2
        model = readCbModel(filename);
    end
end
model.csense(1:size(model.S,1),1)='E';
modelOrig = model;
else
    model=modelOrig;
end

filename =
'Recon3D_301.mat'

```

Set the level of printing, zero for silent, higher for more output.

```
printLevel=2;
```

Choose the directory to place the results

```

basePath='~/work/sbgCloud/';
%resultsPath=[basePath '/programReconstruction/projects/recon2models/results/
reconXs/' model.modelID];
resultsPath=[basePath '/courses/2019_Leiden_COBRA/practicalsDemo/Day4/' 
model.modelID];
resultsFileName=[resultsPath filesep model.modelID];

```

Create and enter the folder for the results if it does not already exist

```

if ~exist(resultsPath,'dir')
    mkdir(resultsPath)
end
cd(resultsPath)

```

Optionally create a diary to save the output in case it is very long, this makes it easier to search, especially when debugging the process during the early stages.

```

if 0
    diary([resultsFileName '_diary.txt'])
end

```

Overview some of the key properties of the reconstruction

Noting the initial size of the reconstruction is useful for comparisons later with subsets derived according to mathematical specifications.

```

[nMet,nRxn]=size(model.S);
fprintf('%6s\t%6s\n','#mets','#rxns')

#mets      #rxns

```

```
fprintf( '%6u\t%6u\t%s\n', nMet, nRxn, ' totals.' )
```

```
8399      13543      totals.
```

Make sure the stoichiometric matrix is stored in a sparse format as this accelerates computations with large networks

```
model.S=sparse(model.S);
```

Check in case the reconstruction is a model that is already ready for flux balance analysis

There is no need to run this live script any further if the reconstruction already satisfies the conditions necessary for flux balance analysis. That is if all internal reactants and reactions are stoichiometrically consistent, and all reactions are flux consistent, then the reconstruction satisfies the criteria to designate it a model ready for flux balance analysis.

SIntMetBool m x 1 Boolean of metabolites heuristically though to be involved in mass balanced reactions.

SIntRxnBool n x 1 Boolean of reactions heuristically though to be mass balanced.

SConsistentMetBool m x 1 Boolean vector indicating consistent mets

SConsistentRxnBool n x 1 Boolean vector indicating consistent rxns

fluxConsistentMetBool m x 1 Boolean vector indicating flux consistent mets

fluxConsistentRxnBool n x 1 Boolean vector indicating flux consistent rxns

```
if all(isfield(model,{ 'SIntMetBool', 'SIntRxnBool', 'SConsistentMetBool', ...
    'SConsistentRxnBool', 'fluxConsistentMetBool', 'fluxConsistentRxnBool' })) ...
    if all(model.SIntMetBool & model.SConsistentMetBool) ...
        && nnz(model.SIntRxnBool &
model.SConsistentRxnBool)==nnz(model.SIntRxnBool) ...
            && all(model.fluxConsistentMetBool) ...
            && all(model.fluxConsistentRxnBool)
        fullyStoichAndFluxConsistent=1;
        fprintf( '%s\n', 'Reconstruction is a model that is already ready for
flux balance analysis')
    end
    return
else
    fullyStoichAndFluxConsistent=0;
    fprintf( '%s\n', 'Reconstruction must be tested to check if it is ready
for flux balance analysis')
end
```

Reconstruction must be tested to check if it is ready for flux balance analysis

Manually remove certain reactions from the reconstruction

Before attempting to algorithmically remove stoichiometrically or flux inconsistent supposed internal reactions from a reconstruction to generate a model, there is an option to review the content of the reconstruction and manually identify reactions for removal. That is, there are two options:

A. Skip manual review of reconstruction content. Move to the next step.

B. Review the content of the reconstruction and omit any reactions that are assumed to be stoichiometrically or flux inconsistent. With respect to stoichiometric inconsistency, such reactions may be obviously mass imbalanced and not satisfy the heuristic conditions for identification as an external reaction. Alternatively, such reactions may be identified by a previous pass through of this tutorial as being of unknown stoichiometric consistent (`model.unknownSConsistencyRxnBool(j)==1`), after the largest stoichiometrically consistent subset of the network has been identified. This is an iterative process where multiple rounds of identification of the largest stoichiometrically consistent set and manual curation of the remainder that is of unknown stoichiometric consistency is necessary.

```
if strcmp(filename, 'Recon3.0model')
    modelOrig=model;
    if 0
        if 1
            %Rename some of the biomass reactions to make them more
            obviously exchange
            %reactions
            model.rxns{strcmp(model.rxns, 'biomass_reaction')}=
            'EX_biomass_reaction';
            model.rxns{strcmp(model.rxns, 'biomass_maintenance')}=
            'EX_biomass_maintenance';
            model.rxns{strcmp(model.rxns, 'biomass_maintenance_noTrTr')}=
            'EX_biomass_maintenance_noTrTr';

            %ATP hydrolysis is not imbalanced like all the other demand
            reactions so
            %give it a different acronym ATPM = ATP Maintenance
            bool=strcmp('DM_atp_c_',model.rxns);
            model.rxns{bool}='ATPM';
        end
        [model,removeMetBool,removeRxnBool] =
        manuallyAdaptRecon3(model,printLevel);
    else
        [model,removeMetBool,removeRxnBool] =
        manuallyAdaptRecon3Ines(model,printLevel);
    end
    [nMet0,nRxn0]=size(modelOrig.S);
    [nMet,nRxn]=size(model.S);
    if nMet0==nMet && nRxn0==nRxn && printLevel>0
        fprintf('%s\n','--- Manually removing rows and columns of the
        stoichiometric matrix----')
        fprintf('%6s\t%6s\n','#mets','#rxns')
```

```

        fprintf('%6u\t%6u\t%s\n',nMet0,nRxn0,' totals.')
        fprintf('%6u\t%6u\t%s\n',nMet0-nMet,nRxn0-nRxn,' manually removed.')
        fprintf('%6u\t%6u\t%s\n',nMet,nRxn,' remaining.')
    end
end

```

Remove any trivial rows and columns of the stoichiometric matrix

Remove any zero rows or columns of the stoichiometric matrix

```

modelOrig=model;
model=removeTrivialStoichiometry(model);
[nMet0,nRxn0]=size(modelOrig.S);
[nMet,nRxn]=size(model.S);
if nMet0==nMet && nRxn0==nRxn && printLevel>0
    fprintf('%s\n','---Checking for Remove any trivial rows and columns of
the stoichiometric matrix----')
    fprintf('%s\t%s\n','#mets','#rxns')
    fprintf('%6u\t%6u\t%s\n',nMet0,nRxn0,' totals.')
    fprintf('%6u\t%6u\t%s\n',nMet0-nMet,nRxn0-nRxn,' duplicates removed.')
    fprintf('%6u\t%6u\t%s\n',nMet,nRxn,' remaining.')
end

---Checking for Remove any trivial rows and columns of the stoichiometric matrix----
#mets      #rxns
8399      13543      totals.
0          0          duplicates removed.
8399      13543      remaining.

```

Check for duplicate columns by detecting the columns of the S matrix that are identical upto scalar multiplication.

```

modelOrig=model;
dupDetectMethod='FR';
dupDetectMethod='S';
removeFlag=0;
[modelOut,removedRxnInd, keptRxnInd] =
checkDuplicateRxn(model,dupDetectMethod,removeFlag,printLevel-2);

```

Remove any duplicate reactions, and uniquely involved reactants, from the stoichiometric matrix.

```

if length(removedRxnInd)>0
    irrevFlag=0;
    metFlag=1;
    %set all reactions reversible that are duplicates
    model.lb(removedRxnInd)=-model.ub(removedRxnInd);
    %remove duplicates
    model = removeRxns(model,model.rxns(removedRxnInd),irrevFlag,metFlag);
end

```

Display the statistics on the duplicate reactions,

```
[nMet0,nRxn0]=size(modelOrig.S);
[nMet,nRxn]=size(model.S);
if nMet0==nMet && nRxn0==nRxn && printLevel>0
    fprintf(' %s\n', '---Remove any duplicate reactions----')
    [nMet0,nRxn0]=size(modelOrig.S);
    [nMet,nRxn]=size(model.S);
    fprintf(' %6s\t%6s\n', '#mets', '#rxns')
    fprintf(' %6u\t%6u\t%6u\n', nMet0,nRxn0, ' totals.')
    fprintf(' %6u\t%6u\t%6u\n', nMet0-nMet, nRxn0-nRxn, ' duplicates removed.')
    fprintf(' %6u\t%6u\t%6u\n', nMet,nRxn, ' remaining.')
end
```

```
--Remove any duplicate reactions---
#mets      #rxns
8399      13543      totals.
0          0          duplicates removed.
8399      13543      remaining.
```

Remove any duplicate reactions upto protons

Remove reactions reactions that differ only in the number of protons involved as substrates or products. Also remove exclusively involved reactants.

Save a temporary model for testing, before making any changes.

```
modelH=model;
```

Find the proton indicies in different compartments. A proton, with index i, is assumed to be represented by an abbreviation within model.mets{i} like h[*], where * denotes the compartment symbol.

```
nMetChars=zeros(length(modelH.mets),1);
for m=1:length(modelH.mets)
    nMetChars(m,1)=length(modelH.mets{m});
end
protonMetBool=strncmp(modelH.mets,'h',1) & nMetChars==length('h[*]');
if printLevel>2
    disp(modelH.mets(protonMetBool))
end
```

Zero out the proton stoichiometric coefficients from the temporary model for testing

```
modelH.S(protonMetBool,:)=0;
```

Check for duplicate columns, upto protons, by detecting the columns of the S matrix that are identical upto scalar multiplication.

```
dupDetectMethod='FR';
removeFlag=0;
```

```
[modelOut, removedRxnInd, keptRxnInd] =
checkDuplicateRxn(modelH, dupDetectMethod, removeFlag, printLevel-1);
```

Checking for reaction duplicates by stoichiometry (up to orientation) ...

Keep: 25HVITD2t 25hvitd2[c] -> 25hvitd2[e]
 Duplicate: 25HVITD2tin 25hvitd2[e] -> 25hvitd2[c]
 Keep: 25HVITD2tin_m 25hvitd2[c] -> 25hvitd2[m]
 Duplicate: 25HVITD2tm 25hvitd2[m] -> 25hvitd2[c]
 Keep: 25HVITD3t 25hvitd3[c] -> 25hvitd3[e]
 Duplicate: 25HVITD3tin 25hvitd3[e] -> 25hvitd3[c]
 Keep: 25HVITD3tin_m 25hvitd3[c] -> 25hvitd3[m]
 Duplicate: 25HVITD3tm 25hvitd3[m] -> 25hvitd3[c]
 Keep: 3MOBt2im 3mob[c] -> 3mob[m]
 Duplicate: HMR_3746 3mob[c] <=> 3mob[m]
 Keep: 5MTHFT 5mthf[e] <=> 5mthf[c]
 Duplicate: MTHFTe 5mthf[c] -> 5mthf[e]
 Keep: ADNt adn[e] <=> adn[c]
 Duplicate: ADNCNT3tc adn[e] <=> adn[c]
 Keep: ADPRIbt adprib[e] -> adprib[c]
 Duplicate: ADPRIbte adprib[c] <=> adprib[e]
 Keep: ALAT4 nal[e] + ala_L[e] -> nal[c] + ala_L[c]
 Duplicate: HMR_9605 nal[e] + ala_L[e] -> nal[c] + ala_L[c]
 Keep: ALCD21_D nad[c] + 12ppd_R[c] -> nadh[c] + lald_D[c]
 Duplicate: PPDOx nadh[c] + lald_D[c] -> nad[c] + 12ppd_R[c]
 Keep: ALCD22_D nad[c] + lald_D[c] -> nadh[c] + mthgxl[c]
 Duplicate: LALDO2x nadh[c] + mthgxl[c] -> nad[c] + lald_D[c]
 Keep: ATPasel h2o[c] + atp[c] -> adp[c] + pi[c]
 Duplicate: DM_atp_c_ h2o[c] + atp[c] -> adp[c] + pi[c]

Warning: BTNt2 has more than one replicate

Keep: BTNt2 btn[e] <=> btn[c]
 Duplicate: BTNTe btn[c] -> btn[e]
 Keep: C14STRr nadph[r] + 44mctr[r] -> nadp[r] + 44mzym[r]
 Duplicate: r0780 nadp[r] + 44mzym[r] <=> nadph[r] + 44mctr[r]
 Keep: C160CPT1 crn[c] + pmtncoa[c] <=> coa[c] + pmtncrn[c]
 Duplicate: C160CPT2rbc coa[c] + pmtncoa[c] <=> crn[c] + pmtncoa[c]
 Keep: C161CPT2 coa[m] + hdcecrn[m] <=> crn[m] + hdcoa[m]
 Duplicate: r0446 crn[m] + hdcoa[m] <=> coa[m] + hdcecrn[m]
 Keep: C181CPT1 crn[c] + odecraa[c] <=> coa[c] + odecraa[c]
 Duplicate: C181CPT2rbc coa[c] + odecraa[c] <=> crn[c] + odecraa[c]
 Keep: CITtam cit[c] + mal_L[m] <=> cit[m] + mal_L[c]
 Duplicate: HMR_4964 cit[c] + mal_L[m] -> cit[m] + mal_L[c]
 Keep: CRNT crn[e] <=> crn[c]
 Duplicate: CRNTHa crn[c] -> crn[e]
 Keep: CRNTuNa nal[e] + crn[e] -> nal[c] + crn[c]
 Duplicate: CRNCT2te nal[c] + crn[c] <=> nal[e] + crn[e]
 Keep: CRVNCTr crvnc[e] <=> crvnc[c]
 Duplicate: CE0328te crvnc[c] <=> crvnc[e]
 Keep: CYST4 nal[e] + cys_L[e] -> nal[c] + cys_L[c]
 Duplicate: CYSSNAT5tc nal[e] + cys_L[e] <=> nal[c] + cys_L[c]
 Keep: CYTdt cytd[e] <=> cytd[c]
 Duplicate: CYTdt2r cytd[e] <=> cytd[c]
 Keep: DALAt2r ala_D[e] <=> ala_D[c]
 Duplicate: ALA-DTDe ala_D[c] -> ala_D[e]
 Keep: DMHPTCRNte dmhptcrn[c] <=> dmhptcrn[e]
 Duplicate: DMHPTCRNtr dmhptcrn[e] <=> dmhptcrn[c]
 Keep: DNDPt10m dadp[c] + dcdp[m] -> dcdp[c] + dadp[m]
 Duplicate: DNDPt29m dcdp[c] + dadp[m] -> dadp[c] + dcdp[m]
 Keep: DNDPt11m dadp[c] + dgdp[m] -> dgdp[c] + dadp[m]
 Duplicate: DNDPt35m dgdp[c] + dadp[m] -> dadp[c] + dgdp[m]
 Keep: DNDPt14m dtdp[m] + dudp[c] -> dt dp[c] + dudp[m]
 Duplicate: DNDPt22m dt dp[c] + dudp[m] -> dt dp[m] + dudp[c]
 Keep: DNDPt15m dgdp[m] + dudp[c] -> dgdp[c] + dudp[m]
 Duplicate: DNDPt33m dgdp[c] + dudp[m] -> dgdp[m] + dudp[c]
 Keep: DNDPt16m dadp[m] + dudp[c] -> dadp[c] + dudp[m]

```

Duplicate: DNDPt8m dadp[c] + dudp[m] -> dadp[m] + dudp[c]
Keep: DNDPt17m dc当地[m] + dudp[c] -> dc当地[c] + dudp[m]
Duplicate: DNDPt26m dc当地[c] + dudp[m] -> dc当地[m] + dudp[c]
Keep: DNDPt23m dgdp[m] + dtdp[c] -> dgdp[c] + dtdp[m]
Duplicate: DNDPt34m dgdp[c] + dtdp[m] -> dgdp[m] + dtdp[c]
Keep: DNDPt24m dadp[m] + dtdp[c] -> dadp[c] + dtdp[m]
Duplicate: DNDPt9m dadp[c] + dtdp[m] -> dadp[m] + dtdp[c]
Keep: DNDPt25m dc当地[m] + dtdp[c] -> dc当地[c] + dtdp[m]
Duplicate: DNDPt27m dc当地[c] + dtdp[m] -> dc当地[m] + dtdp[c]
Keep: DNDPt28m dc当地[c] + dgdp[m] -> dgdp[c] + dc当地[m]
Duplicate: DNDPt36m dgdp[c] + dc当地[m] -> dc当地[c] + dgdp[m]
Keep: DOPAtu dopa[e] <=> dopa[c]
Duplicate: DOPAENT4tc dopa[e] <=> dopa[c]
Keep: EBP2r zymstnl[r] -> lthstrl[r]
Duplicate: r1381 lthstrl[r] <=> zymstnl[r]
Keep: FE2t fe2[e] -> fe2[c]
Duplicate: FE2DMT1 fe2[e] -> fe2[c]
Keep: FE2tm fe2[c] -> fe2[m]
Duplicate: HMR_5420 fe2[c] -> fe2[m]
Keep: FUCFUCFUCGALACGLC13GALACGLCGAL14ACGLCGALGLUSIDe fucfucfucgalacglc13galacglcgall4acglcg
Duplicate: HMR_9651 fucfucfucgalacglc13galacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Keep: FUCFUCFUCGALACGLCGAL14ACGLCGALGLUSIDe fucfucfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcg
Duplicate: HMR_9645 fucfucfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Keep: FUCGALFUCGALACGLCGALGLUSIDe fucgalfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Duplicate: HMR_9643 fucgalfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Keep: GALFUCGALACGLCGAL14ACGLCGALGLUSIDe galfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Duplicate: HMR_9646 galfucgalacglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcgall4acglcg
Keep: GALt1r gal[e] <=> gal[c]
Duplicate: GALt2_2 gal[e] <=> gal[c]
Keep: GDPtg gdp[c] <=> gdp[g]
Duplicate: HMR_7743 gdp[c] <=> gdp[g]

Warning: GLCt1r has more than one replicate
Keep: GLCt1r glc_D[e] <=> glc_D[c]
Duplicate: GLCGLUT2 glc_D[c] -> glc_D[e]
Keep: GLNtm gln_L[c] -> gln_L[m]
Duplicate: HMR_5101 gln_L[c] -> gln_L[m]
Keep: GLYC3Ptm glyc3p[c] -> glyc3p[m]
Duplicate: GLYC3Ptmc glyc3p[m] <=> glyc3p[c]
Keep: GLYT4 na1[e] + gly[e] -> na1[c] + gly[c]
Duplicate: GLYSNAT5tc na1[e] + gly[e] <=> na1[c] + gly[c]
Keep: GSNT gsn[e] <=> gsn[c]
Duplicate: GSNT2r gsn[e] <=> gsn[c]
Keep: HIST4 na1[e] + his_L[e] -> na1[c] + his_L[c]
Duplicate: HISSNAT5tc na1[e] + his_L[e] <=> na1[c] + his_L[c]
Keep: HISTiDF his_L[e] -> his_L[c]
Duplicate: HISCAT1 his_L[c] <=> his_L[e]
Keep: HSD17B7r nadph[r] + estrone[r] -> nadp[r] + estradiol[r]
Duplicate: HMR_2041 nadph[r] + estrone[r] -> nadp[r] + estradiol[r]

Warning: Htg has more than one replicate
Keep: Htg <=>
Duplicate: Htmi ->
Keep: INST ins[e] <=> ins[c]
Duplicate: INST2 ins[e] <=> ins[c]
Keep: L_LACTcm lac_L[c] -> lac_L[m]
Duplicate: L_LACTm lac_L[c] -> lac_L[m]
Keep: LNLCPT1 crn[c] + lnlc当地[c] <=> coa[c] + lnlc当地[c]
Duplicate: LNLCPT2rbc coa[c] + lnlc当地[c] <=> crn[c] + lnlc当地[c]

Warning: NACUP has more than one replicate
Keep: NACUP nac[e] -> nac[c]
Duplicate: NACHORCTL3le nac[e] -> nac[c]
Keep: NADHtpu nadh[c] -> nadh[x]
Duplicate: NADtpu nadh[x] -> nadh[c]
Keep: NAT na1[e] <=> na1[c]
Duplicate: NAT3_1 na1[c] <=> na1[e]

```

```

    Keep: NCAMUP ncam[e] -> ncam[c]
Duplicate: NCAMDe ncam[c] -> ncam[e]
    Keep: NH4t3r nh4[c] <=> nh4[e]
Duplicate: NH4tb nh4[e] <=> nh4[c]
    Keep: NOT no[e] <=> no[c]
Duplicate: NODE no[c] <=> no[e]
    Keep: OCTAt octa[e] <=> octa[c]
Duplicate: OCTAtc octa[c] <=> octa[e]
Warning: ORNt4m has more than one replicate
    Keep: ORNt4m orn[m] + citr_L[c] <=> orn[c] + citr_L[m]
Duplicate: r2412 orn[c] + citr_L[m] -> orn[m] + citr_L[c]
    Keep: P5CRxm nadh[m] + 1pyr5c[m] -> nadh[m] + pro_L[m]
Duplicate: PRO1xm nadh[m] + pro_L[m] -> nadh[m] + 1pyr5c[m]
    Keep: PITx pi[c] <=> pi[x]
Duplicate: HMR_5344 pi[c] <=> pi[x]
    Keep: PRODt2r pro_D[e] <=> pro_D[c]
Duplicate: PRO_Dtde pro_D[c] <=> pro_D[e]
    Keep: RIBt rib_D[e] <=> rib_D[c]
Duplicate: RIBT2 rib_D[e] -> rib_D[c]
    Keep: SRTNTu srtn[e] <=> srtn[c]
Duplicate: SRTNENT4tc srtn[e] <=> srtn[c]
    Keep: SUCCtp succ[c] <=> succ[x]
Duplicate: SUCCTD succ[x] <=> succ[c]
    Keep: TAGt tag_hs[e] <=> tag_hs[c]
Duplicate: TAGHSTDe tag_hs[c] -> tag_hs[e]
Warning: THYMDt1 has more than one replicate
    Keep: THYMDt1 thymd[e] -> thymd[c]
Duplicate: THMDt2r thymd[e] <=> thymd[c]
    Keep: TRDRm nadph[m] + trdox[m] -> nadp[m] + trdrd[m]
Duplicate: r1433 nadp[m] + trdrd[m] -> nadph[m] + trdox[m]
    Keep: URIt uri[e] <=> uri[c]
Duplicate: URIt2r uri[e] <=> uri[c]
    Keep: VITD3t vitd3[c] -> vitd3[e]
Duplicate: VITD3t2 vitd3[e] -> vitd3[c]
Warning: VITD3tm has more than one replicate
    Keep: VITD3tm vitd3[m] -> vitd3[c]
Duplicate: HMR_2116 vitd3[c] <=> vitd3[m]
    Keep: XOLEST2te xolest2_hs[e] <=> xolest2_hs[c]
Duplicate: XOLEST2HSTDle xolest2_hs[c] -> xolest2_hs[e]
    Keep: r0276 nh4[c] + nadp[c] + imp[c] <=> nadph[c] + gmp[c]
Duplicate: GMPR nadph[c] + gmp[c] -> nh4[c] + nadp[c] + imp[c]
    Keep: r0488 2 nadp[c] + coa[c] + mev_R[c] <=> 2 nadph[c] + hmgcoa[c]
Duplicate: HMGCOARc 2 nadph[c] + hmgcoa[c] -> 2 nadp[c] + coa[c] + mev_R[c]
    Keep: r0537 ethamp[c] + hxdcal[c] -> sph1p[c]
Duplicate: SGPL11c sph1p[c] -> ethamp[c] + hxdcal[c]
    Keep: r0561 coa[m] + 2mpdhl[m] -> ibcoa[m] + dhlam[m]
Duplicate: RE3326M ibcoa[m] + dhlam[m] <=> coa[m] + 2mpdhl[m]
    Keep: r0808 HC00004[c] -> HC00004[e]
Duplicate: HC00004t1e HC00004[e] -> HC00004[c]
    Keep: r0817 citr_L[c] <=> citr_L[e]
Duplicate: CITRtr citr_L[e] <=> citr_L[c]
    Keep: r0839 orot[e] <=> orot[c]
Duplicate: OROte orot[e] -> orot[c]
    Keep: r0899 ala_B[c] <=> ala_B[e]
Duplicate: BALAPAT1tc ala_B[e] -> ala_B[c]
    Keep: r0913 icit[m] + mal_L[c] <=> mal_L[m] + icit[c]
Duplicate: r2387 mal_L[m] + icit[c] -> icit[m] + mal_L[c]
    Keep: r0915 cit[c] + succ[m] <=> cit[m] + succ[c]
Duplicate: r2382 cit[c] + succ[m] -> cit[m] + succ[c]
    Keep: r0944 spmd[c] <=> spmd[e]
Duplicate: SPMTDe spmd[e] <=> spmd[c]
    Keep: r1050 chsterol[e] <=> chsterol[c]
Duplicate: CHOLESTTTDe chsterol[c] -> chsterol[e]
    Keep: r1067 his_L[l] -> his_L[c]

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Duplicate: HIShPTtc his_L[1] -> his_L[c]
Keep: r1078 tyr_L[c] -> tyr_L[m]
Duplicate: HMR_5099 tyr_L[c] <=> tyr_L[m]
Keep: r1127 HC00005[c] -> HC00005[r]
Duplicate: HC00005t1r HC00005[r] -> HC00005[c]
Keep: r1128 HC00009[c] -> HC00009[r]
Duplicate: HC00009t1r HC00009[r] -> HC00009[c]
Keep: r1129 HC00004[c] -> HC00004[r]
Duplicate: HC00004t1r HC00004[r] -> HC00004[c]
Keep: r1131 HC00006[c] -> HC00006[r]
Duplicate: HC00006t1r HC00006[r] -> HC00006[c]
Keep: r1132 HC00007[c] -> HC00007[r]
Duplicate: HC00007t1r HC00007[r] -> HC00007[c]
Keep: r1133 HC00008[c] -> HC00008[r]
Duplicate: HC00008t1r HC00008[r] -> HC00008[c]
Keep: r1147 akg[c] + icit[m] <=> akg[m] + icit[c]
Duplicate: r2385 akg[m] + icit[c] -> akg[c] + icit[m]
Keep: r1155 2obut[c] -> 2obut[m]
Duplicate: r1454 2obut[m] -> 2obut[c]
Keep: r1423 pi[c] -> pi[e]
Duplicate: PI6b pi[e] <=> pi[c]
Keep: r1427 his_L[c] -> his_L[m]
Duplicate: r2416 his_L[m] -> his_L[c]
Keep: r1429 glyc3p[c] <=> glyc3p[x]
Duplicate: GLY3Pt glyc3p[x] -> glyc3p[c]
Keep: r1441 trdrd[c] -> trdrd[m]
Duplicate: HMR_6618 trdrd[c] <=> trdrd[m]
Keep: r1455 phe_L[c] -> phe_L[m]
Duplicate: r1456 phe_L[m] -> phe_L[c]
Keep: r1618 tyr_L[c] + phe_L[e] <=> phe_L[c] + tyr_L[e]
Duplicate: TYRPHELAT2tc phe_L[c] + tyr_L[e] -> tyr_L[c] + phe_L[e]
Keep: r1619 cys_L[c] + phe_L[e] <=> cys_L[e] + phe_L[c]
Duplicate: CYSPHELAT2tc cys_L[e] + phe_L[c] -> cys_L[c] + phe_L[e]
Keep: r1620 leu_L[c] + phe_L[e] <=> leu_L[e] + phe_L[c]
Duplicate: LEUPHELAT2tc leu_L[e] + phe_L[c] <=> leu_L[c] + phe_L[e]
Keep: r1622 asn_L[c] + phe_L[e] <=> asn_L[e] + phe_L[c]
Duplicate: ASNPHELAT2tc asn_L[e] + phe_L[c] -> asn_L[c] + phe_L[e]
Keep: r1623 phe_L[e] + val_L[c] <=> phe_L[c] + val_L[e]
Duplicate: VALPHELAT2tc phe_L[c] + val_L[e] -> phe_L[e] + val_L[c]
Keep: r1624 thr_L[c] + phe_L[e] <=> thr_L[e] + phe_L[c]
Duplicate: THRPHELAT2tc thr_L[e] + phe_L[c] -> thr_L[c] + phe_L[e]
Keep: r1626 ile_L[c] + phe_L[e] <=> ile_L[e] + phe_L[c]
Duplicate: ILEPHELAT2tc ile_L[e] + phe_L[c] -> ile_L[c] + phe_L[e]
Keep: r1644 leu_L[e] + val_L[c] <=> leu_L[c] + val_L[e]
Duplicate: VALLAT1tc leu_L[c] + val_L[e] -> leu_L[e] + val_L[c]
Keep: r1647 ile_L[c] + leu_L[e] <=> ile_L[e] + leu_L[c]
Duplicate: ILELAT1tc ile_L[e] + leu_L[c] -> ile_L[c] + leu_L[e]
Keep: r1668 arg_L[e] + his_L[c] <=> arg_L[c] + his_L[e]
Duplicate: HISyLATthc arg_L[c] + his_L[e] -> arg_L[e] + his_L[c]
Keep: r2009 ala_L[c] + arg_L[e] -> ala_L[e] + arg_L[c]
Duplicate: ALAyLATthc ala_L[e] + arg_L[c] -> ala_L[c] + arg_L[e]
Keep: r2010 gln_L[c] + arg_L[e] -> gln_L[e] + arg_L[c]
Duplicate: GLNyLATthc gln_L[e] + arg_L[c] -> gln_L[c] + arg_L[e]
Keep: r2012 arg_L[e] + met_L[c] -> arg_L[c] + met_L[e]
Duplicate: METyLATthc arg_L[c] + met_L[e] -> arg_L[e] + met_L[c]
Keep: r2014 arg_L[e] + phe_L[c] -> arg_L[c] + phe_L[e]
Duplicate: PHEyLATthc arg_L[c] + phe_L[e] -> arg_L[e] + phe_L[c]
Keep: r2017 arg_L[e] + leu_L[c] -> arg_L[c] + leu_L[e]
Duplicate: LEUyLAThtc arg_L[c] + leu_L[e] -> arg_L[e] + leu_L[c]
Keep: r2073 zn2[e] -> zn2[c]
Duplicate: r2465 zn2[c] -> zn2[e]
Keep: r2346 wharachd[e] <=> wharachd[c]
Duplicate: WHARACHDtd wharachd[c] <=> wharachd[e]
Keep: r2355 HC02203[e] <=> HC02203[c]

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Duplicate: C05953td HC02203[c] <=> HC02203[e]
Keep: r2364 HC02213[e] <=> HC02213[c]
Duplicate: C06439td HC02213[c] <=> HC02213[e]
Keep: r2373 akg[c] + cit[m] <=> akg[m] + cit[c]
Duplicate: r2381 akg[m] + cit[c] -> akg[c] + cit[m]
Keep: r2374 cit[m] + oxa[c] <=> cit[c] + oxa[m]
Duplicate: r2384 cit[c] + oxa[m] -> cit[m] + oxa[c]
Keep: r2375 icit[m] + succ[c] <=> icit[c] + succ[m]
Duplicate: r2386 icit[c] + succ[m] -> icit[m] + succ[c]
Keep: r2376 icit[m] + oxa[c] <=> icit[c] + oxa[m]
Duplicate: r2388 icit[c] + oxa[m] -> icit[m] + oxa[c]
Keep: r2377 akg[c] + HC00342[m] <=> akg[m] + HC00342[c]
Duplicate: r2389 akg[m] + HC00342[c] -> akg[c] + HC00342[m]
Keep: r2378 succ[c] + HC00342[m] <=> succ[m] + HC00342[c]
Duplicate: r2390 succ[m] + HC00342[c] -> succ[c] + HC00342[m]
Keep: r2379 mal_L[c] + HC00342[m] <=> mal_L[m] + HC00342[c]
Duplicate: r2391 mal_L[m] + HC00342[c] -> mal_L[c] + HC00342[m]
Keep: r2380 oxa[c] + HC00342[m] <=> HC00342[c] + oxa[m]
Duplicate: r2392 HC00342[c] + oxa[m] -> oxa[c] + HC00342[m]
Keep: r2395 lys_L[m] + arg_L[c] -> lys_L[c] + arg_L[m]
Duplicate: r2399 lys_L[c] + arg_L[m] -> lys_L[m] + arg_L[c]
Keep: r2396 orn[c] + lys_L[m] -> lys_L[c] + orn[m]
Duplicate: r2403 lys_L[c] + orn[m] -> orn[c] + lys_L[m]
Keep: r2397 lys_L[m] + his_L[c] -> lys_L[c] + his_L[m]
Duplicate: r2406 lys_L[c] + his_L[m] -> lys_L[m] + his_L[c]
Keep: r2398 lys_L[m] + citr_L[c] -> lys_L[c] + citr_L[m]
Duplicate: r2410 lys_L[c] + citr_L[m] -> lys_L[m] + citr_L[c]
Keep: r2400 orn[c] + arg_L[m] -> arg_L[c] + orn[m]
Duplicate: r2404 arg_L[c] + orn[m] -> orn[c] + arg_L[m]
Keep: r2401 arg_L[m] + his_L[c] -> arg_L[c] + his_L[m]
Duplicate: r2407 arg_L[c] + his_L[m] -> arg_L[m] + his_L[c]
Keep: r2402 arg_L[m] + citr_L[c] -> arg_L[c] + citr_L[m]
Duplicate: r2411 arg_L[c] + citr_L[m] -> arg_L[m] + citr_L[c]
Keep: r2405 orn[m] + his_L[c] -> orn[c] + his_L[m]
Duplicate: r2408 orn[c] + his_L[m] -> orn[m] + his_L[c]
Keep: r2409 citr_L[c] + his_L[m] -> citr_L[m] + his_L[c]
Duplicate: r2413 citr_L[m] + his_L[c] -> citr_L[c] + his_L[m]
Keep: r2471 ser_L[e] -> ser_L[c]
Duplicate: r2526 ser_L[e] <=> ser_L[c]
Keep: r2516 lac_L[x] <=> lac_L[c]
Duplicate: LACLt lac_L[x] -> lac_L[c]
Keep: RE3628M dc2coa[m] <=> dece3coa[m]
Duplicate: FAOXC101m dece3coa[m] -> dc2coa[m]
Keep: BCRNe 3bcrn[c] -> 3bcrn[e]
Duplicate: 3BCRNtr 3bcrn[e] <=> 3bcrn[c]
Keep: C101CRNe c101crn[c] -> c101crn[e]
Duplicate: C101CRNtr c101crn[e] <=> c101crn[c]
Keep: C10CRNe c10crn[c] -> c10crn[e]
Duplicate: C10CRNtr c10crn[e] <=> c10crn[c]
Keep: C10DCe c10dc[c] -> c10dc[e]
Duplicate: C10DCtr c10dc[e] <=> c10dc[c]
Keep: C12DCe c12dc[c] -> c12dc[e]
Duplicate: C12DCtr c12dc[e] <=> c12dc[c]
Keep: C141OHe 3tetd7ecoacrn[c] -> 3tetd7ecoacrn[e]
Duplicate: 3TETD7ECOACRNtr 3tetd7ecoacrn[e] <=> 3tetd7ecoacrn[c]
Keep: C142OHe 3ttetddcoacrn[c] -> 3ttetddcoacrn[e]
Duplicate: 3TTETDDCOACRNtr 3ttetddcoacrn[e] <=> 3ttetddcoacrn[c]
Keep: C162OHe 3thexddcoacrn[c] -> 3thexddcoacrn[e]
Duplicate: 3THEXDDCOACRNtr 3thexddcoacrn[e] <=> 3thexddcoacrn[c]
Keep: C16DCe c16dc[c] -> c16dc[e]
Duplicate: C16DCtr c16dc[e] <=> c16dc[c]
Keep: C3DCe c3dc[c] -> c3dc[e]
Duplicate: C3DCtr c3dc[e] <=> c3dc[c]
Keep: C4CRNe c4crn[c] -> c4crn[e]

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Duplicate: C4CRNtr    c4crn[e]      <=>    c4crn[c]
Keep:   C4DCe       c4dc[c]       ->    c4dc[e]
Duplicate: C4DCtr     c4dc[e]      <=>    c4dc[c]
Keep:   C5DCe       c5dc[c]       ->    c5dc[e]
Duplicate: C5DCtr     c5dc[e]      <=>    c5dc[c]
Keep:   C6CRNe      c6crn[c]      ->    c6crn[e]
Duplicate: C6CRNtr    c6crn[e]      <=>    c6crn[c]
Keep:   C6DCe       c6dc[c]       ->    c6dc[e]
Duplicate: C6DCtr     c6dc[e]      <=>    c6dc[c]
Keep:   C81CRNe     c81crn[c]     ->    c81crn[e]
Duplicate: C81CRNtr   c81crn[e]     <=>    c81crn[c]
Keep:   C8CRNe      c8crn[c]      ->    c8crn[e]
Duplicate: C8CRNtr    c8crn[e]      <=>    c8crn[c]
Keep:   C8DCe       c8dc[c]       ->    c8dc[e]
Duplicate: C8DCtr     c8dc[e]      <=>    c8dc[c]
Keep:   DDCRNe      3ddcrn[c]     ->    3ddcrn[e]
Duplicate: 3DDCRNtr   3ddcrn[e]     <=>    3ddcrn[c]
Keep:   DDECCRNe    ddeccrn[c]     ->    ddeccrn[e]
Duplicate: DDECCRNtr  ddeccrn[e]     <=>    ddeccrn[c]
Keep:   DDECE1CRNe   ddecelcrn[c]    ->    ddecelcrn[e]
Duplicate: DDECE1CRNtr  ddecelcrn[e]    <=>    ddecelcrn[c]
Keep:   DECCRNNe    3deccrn[c]     ->    3deccrn[e]
Duplicate: 3DECCRNtr  3deccrn[e]     <=>    3deccrn[c]
Keep:   DECDICRNNe  decdicrn[c]     ->    decdicrn[e]
Duplicate: DECDICRNtr  decdicrn[e]     <=>    decdicrn[c]
Keep:   HEDCECRNe   3hdececrn[c]    ->    3hdececrn[e]
Duplicate: 3HDECECRNtr  3hdececrn[e]    <=>    3hdececrn[c]
Keep:   HEXDCRNe    3hexdcrn[c]    ->    3hexdcrn[e]
Duplicate: 3HEXDCRNtr  3hexdcrn[e]    <=>    3hexdcrn[c]
Keep:   HIVCRNe     3ivcrn[c]      ->    3ivcrn[e]
Duplicate: 3IVCRNtr   3ivcrn[e]      <=>    3ivcrn[c]
Keep:   HOCTDEC2CRNe  3octdec2crn[c]   ->    3octdec2crn[e]
Duplicate: 3OCTDEC2CRNtr  3octdec2crn[e]   <=>    3octdec2crn[c]
Keep:   HOCTDECCRNNe  3octdeccrn[c]    ->    3octdeccrn[e]
Duplicate: 3OCTDECCRNtr  3octdeccrn[e]    <=>    3octdeccrn[c]
Keep:   HTDCRNe     3tdcrn[c]      ->    3tdcrn[e]
Duplicate: 3TDCRNtr   3tdcrn[e]      <=>    3tdcrn[c]
Keep:   IVCRNNe     ivcrn[c]       ->    ivcrn[e]
Duplicate: IVCRNtr    ivcrn[e]      <=>    ivcrn[c]
Keep:   OCTDECE1CRNe  3octdecelcrn[c]   ->    3octdecelcrn[e]
Duplicate: 3OCTDECE1CRNtr  3octdecelcrn[e]   <=>    3octdecelcrn[c]
Keep:   TDCRNe      ttdcrn[c]      ->    ttdcrn[e]
Duplicate: TTDCRNtr   ttdcrn[e]      <=>    ttdcrn[c]
Keep:   TETDEC2CRNe   tetdec2crn[c]    ->    tetdec2crn[e]
Duplicate: TETDEC2CRNtr  tetdec2crn[e]    <=>    tetdec2crn[c]
Keep:   TETDECE1CRNe  tetdecelcrn[c]   ->    tetdecelcrn[e]
Duplicate: TETDECE1CRNtr  tetdecelcrn[e]   <=>    tetdecelcrn[c]
Keep:   TIGCRNNe    c51crn[c]      ->    c51crn[e]
Duplicate: C51CRNtr   c51crn[e]      <=>    c51crn[c]
Keep:   CARPEPT1tc   carn[e]       ->    carn[c]
Duplicate: CARNtr     carn[e]       <=>    carn[c]
Keep:   CBLTDe      adocbl[c]     ->    adocbl[e]
Duplicate: CBLtle     adocbl[e]     ->    adocbl[c]
Keep:   FOLTle      fol[e]       ->    fol[c]
Duplicate: r0963      fol[e]       ->    fol[c]
Keep:   GLYPROPEPT1tc  glypro[e]    ->    glypro[c]
Duplicate: GLYPROT     glypro[c]     <=>    glypro[e]
Keep:   LEULEUPEPT1tc  leuleu[e]    ->    leuleu[c]
Duplicate: LEULEUT     leuleu[c]     <=>    leuleu[e]
Keep:   PNTORDe     pnto_R[c]    ->    pnto_R[e]
Duplicate: PNTOte      pnto_R[e]    <=>    pnto_R[c]
Keep:   PROGLYPEPT1tc  progly[e]   ->    progly[c]
Duplicate: PROGLyt     progly[c]    <=>    progly[e]
Keep:   SBTle       sbt_D[e]     ->    sbt_D[c]

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Duplicate: SBT_Dtde sbt_D[c] <=> sbt_D[e]
Keep: TAUPAT1c taur[e] -> taur[c]
Duplicate: TAURCHAE taur[c] -> taur[e]
Keep: GLYCTDle glyc[e] <=> glyc[c]
Duplicate: GLYCT glyc[c] <=> glyc[e]
Keep: KHte k[e] <=> k[c]
Duplicate: r1492 k[c] -> k[e]
Keep: PHEMEE pheme[c] -> pheme[e]
Duplicate: PHEMET pheme[e] -> pheme[c]
Keep: SPRMTDe sprm[e] <=> sprm[c]
Duplicate: SPRMt2r sprm[e] <=> sprm[c]
Keep: BALABETAtc2 cala[e] <=> cala[c]
Duplicate: CALAtr cala[e] <=> cala[c]
Keep: CRTNtr crtn[e] <=> crtn[c]
Duplicate: HMR_9619 crtn[e] -> crtn[c]
Keep: ALAPAT4te ala_L[e] <=> ala_L[c]
Duplicate: ALAT2r ala_L[e] <=> ala_L[c]
Keep: PROPAT4te pro_L[e] <=> pro_L[c]
Duplicate: PROT2r pro_L[e] <=> pro_L[c]
Keep: 5AOPT 5aop[c] <=> 5aop[e]
Duplicate: 5AOPT2 5aop[e] -> 5aop[c]
Keep: ABT_Dt abt_D[e] <=> abt_D[c]
Duplicate: ABT_Dt2 abt_D[e] <=> abt_D[c]
Keep: ELAIDCRNtd elaidcrn[c] <=> elaidcrn[e]
Duplicate: ELAIDCRNtr elaidcrn[e] <=> elaidcrn[c]
Keep: HC02149td pcrn[c] <=> pcrn[e]
Duplicate: PCRNtr pcrn[e] <=> pcrn[c]
Keep: LNLCCRNTd lnlccrn[c] <=> lnlccrn[e]
Duplicate: LNLCCRNRtr lnlccrn[e] <=> lnlccrn[c]
Keep: PCSsec pcs[c] -> pcs[e]
Duplicate: PCSup pcs[e] -> pcs[c]
Keep: 3HCINNMup 3hcinnm[e] -> 3hcinnm[c]
Duplicate: 3HCINNMsec 3hcinnm[c] -> 3hcinnm[e]
Keep: 3HPPAUp 3hppa[e] -> 3hppa[c]
Duplicate: 3HPPAsec 3hppa[c] -> 3hppa[e]
Keep: PACALDtm pacald[c] <=> pacald[m]
Duplicate: HMR_4684 pacald[c] <=> pacald[m]
Keep: ACNAMt2 acnam[e] -> acnam[c]
Duplicate: ACNAMtr acnam[c] -> acnam[e]
Keep: ETHAt etha[e] <=> etha[c]
Duplicate: ETHAtr etha[c] -> etha[e]
Keep: THMtrbc thm[e] <=> thm[c]
Duplicate: THMT3 thm[e] <=> thm[c]
Keep: BUTT2r but[e] <=> but[c]
Duplicate: HMR_0155 but[e] <=> but[c]
Keep: DIGALSGALSIDECt digalsgalside_hs[c] -> digalsgalside_hs[e]
Duplicate: DIGALSGALSIDEt1 digalsgalside_hs[e] -> digalsgalside_hs[c]
Keep: PAIL_hs_SECt pail_hs[c] -> pail_hs[e]
Duplicate: PAIL_hs_t1e pail_hs[e] -> pail_hs[c]
Keep: PAILPALM_HSSECT pailpalm_hs[c] -> pailpalm_hs[e]
Duplicate: PAILPALM_HSt1e pailpalm_hs[e] -> pailpalm_hs[c]
Keep: PAILR_HSSECT pailar_hs[c] -> pailar_hs[e]
Duplicate: PAILR_HSt1e pailar_hs[e] -> pailar_hs[c]
Keep: PAILSTE_HSSECT pailste_hs[c] -> pailste_hs[e]
Duplicate: PAILSTE_HSt1e pailste_hs[e] -> pailste_hs[c]
Keep: SPHMYLN180241_hs_SECt sphmyln180241_hs[c] -> sphmyln180241_hs[e]
Duplicate: SPHMYLN180241_hs_t1 sphmyln180241_hs[e] -> sphmyln180241_hs[c]
Keep: SPHMYLN18114_hs_SECt sphmyln18114_hs[c] -> sphmyln18114_hs[e]
Duplicate: SPHMYLN18114_hs_t1 sphmyln18114_hs[e] -> sphmyln18114_hs[c]
Keep: SPHMYLN18115_hs_SECt sphmyln18115_hs[c] -> sphmyln18115_hs[e]
Duplicate: SPHMYLN18115_hs_t1 sphmyln18115_hs[e] -> sphmyln18115_hs[c]
Keep: SPHMYLN18116_hs_SECt sphmyln18116_hs[c] -> sphmyln18116_hs[e]
Duplicate: SPHMYLN18116_hs_t1 sphmyln18116_hs[e] -> sphmyln18116_hs[c]
Keep: SPHMYLN181161_hs_SECt sphmyln181161_hs[c] -> sphmyln181161_hs[e]

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Duplicate: SPHMYLN181161_hs_t1 sphmyln181161_hs[e] -> sphmyln181161_hs[c]
Keep: SPHMYLN18117_hs_SECt sphmyln18117_hs[c] -> sphmyln18117_hs[e]
Duplicate: SPHMYLN18117_hs_t1 sphmyln18117_hs[e] -> sphmyln18117_hs[c]
Keep: SPHMYLN18118_hs_SECt sphmyln18118_hs[c] -> sphmyln18118_hs[e]
Duplicate: SPHMYLN18118_hs_t1 sphmyln18118_hs[e] -> sphmyln18118_hs[c]
Keep: SPHMYLN181181_hs_SECt sphmyln181181_hs[c] -> sphmyln181181_hs[e]
Duplicate: SPHMYLN181181_hs_t1 sphmyln181181_hs[e] -> sphmyln181181_hs[c]
Keep: SPHMYLN18120_hs_SECt sphmyln18120_hs[c] -> sphmyln18120_hs[e]
Duplicate: SPHMYLN18120_hs_t1 sphmyln18120_hs[e] -> sphmyln18120_hs[c]
Keep: SPHMYLN181201_hs_SECt sphmyln181201_hs[c] -> sphmyln181201_hs[e]
Duplicate: SPHMYLN181201_hs_t1 sphmyln181201_hs[e] -> sphmyln181201_hs[c]
Keep: SPHMYLN18121_hs_SECt sphmyln18121_hs[c] -> sphmyln18121_hs[e]
Duplicate: SPHMYLN18121_hs_t1 sphmyln18121_hs[e] -> sphmyln18121_hs[c]
Keep: SPHMYLN18122_hs_SECt sphmyln18122_hs[c] -> sphmyln18122_hs[e]
Duplicate: SPHMYLN18122_hs_t1 sphmyln18122_hs[e] -> sphmyln18122_hs[c]
Keep: SPHMYLN181221_hs_SECt sphmyln181221_hs[c] -> sphmyln181221_hs[e]
Duplicate: SPHMYLN181221_hs_t1 sphmyln181221_hs[e] -> sphmyln181221_hs[c]
Keep: SPHMYLN18123_hs_SECt sphmyln18123_hs[c] -> sphmyln18123_hs[e]
Duplicate: SPHMYLN18123_hs_t1 sphmyln18123_hs[e] -> sphmyln18123_hs[c]
Keep: SPHMYLN1824_hs_SECt sphmyln1824_hs[c] -> sphmyln1824_hs[e]
Duplicate: SPHMYLN1824_hs_t1 sphmyln1824_hs[e] -> sphmyln1824_hs[c]
Keep: SPHMYLN1825_hs_SECt sphmyln1825_hs[c] -> sphmyln1825_hs[e]
Duplicate: SPHMYLN1825_hs_t1 sphmyln1825_hs[e] -> sphmyln1825_hs[c]
Keep: 3AIBt1 3aib[e] <=> 3aib[c]
Duplicate: HMR_8090 3aib[c] -> 3aib[e]
Keep: 2HXIC_Lt1e 2hxic_L[e] -> 2hxic_L[c]
Duplicate: 2HXIC_Lt2e 2hxic_L[c] -> 2hxic_L[e]
Keep: MMAt2e mma[c] <=> mma[e]
Duplicate: MMAtte mma[e] <=> mma[c]
Keep: CE4890te2 CE4890[c] <=> CE4890[e]
Duplicate: CE4890te CE4890[c] <=> CE4890[e]
Keep: MLTHFTe mlthf[e] -> mlthf[c]
Duplicate: MLTHFTe3 mlthf[e] -> mlthf[c]
Keep: TYMte2 tym[c] <=> tym[e]
Duplicate: TYMte tym[c] <=> tym[e]
Keep: 1A25DHVITD3te 1a25dhvitzd3[e] -> 1a25dhvitzd3[c]
Duplicate: 1A25DHVITD3t2e 1a25dhvitzd3[c] -> 1a25dhvitzd3[e]
Keep: ORN_Dtx orn_D[x] <=> orn_D[c]
Duplicate: HMR_9179 orn_D[c] <=> orn_D[x]
Keep: ORN_Dte orn_D[c] <=> orn_D[e]
Duplicate: HMR_9180 orn_D[c] <=> orn_D[e]
Keep: HC00005te HC00005[c] -> HC00005[e]
Duplicate: HC00005t1e HC00005[e] -> HC00005[c]
Keep: HC00006te HC00006[c] -> HC00006[e]
Duplicate: HC00006t1e HC00006[e] -> HC00006[c]
Keep: HC00007te HC00007[c] -> HC00007[e]
Duplicate: HC00007t1e HC00007[e] -> HC00007[c]
Keep: HC00008te HC00008[c] -> HC00008[e]
Duplicate: HC00008t1e HC00008[e] -> HC00008[c]
Keep: HC00009te HC00009[c] -> HC00009[e]
Duplicate: HC00009t1e HC00009[e] -> HC00009[c]
Keep: NO2te no2[e] <=> no2[c]
Duplicate: HMR_6991 no2[c] <=> no2[e]
Keep: HMR_0025 M01268[n] -> M01268[c]
Duplicate: HMR_0030 M01268[c] -> M01268[n]
Keep: HMR_9581 M02035[c] <=> M02035[e]
Duplicate: HMR_9582 M02035[e] -> M02035[c]
Keep: HMR_9583 M02467[c] <=> M02467[e]
Duplicate: HMR_9584 M02467[e] -> M02467[c]
Keep: HMR_0031 0.0024 ak2gchol_hs[c] + 0.0008 dak2gpe_hs[c] + 0.0016 pail_hs[c] + 0.19 dag_hs[c]
Duplicate: HMR_0032 M02392[c] -> 0.0024 ak2gchol_hs[c] + 0.0008 dak2gpe_hs[c] + 0.0016 pail_hs[c]
Keep: ALLOP2tu allop[e] -> allop[c]
Duplicate: ALLOPtepb allop[e] <=> allop[c]
Keep: ATVCACIDMCTtu atvacid[e] <=> atvacid[c]

```

```

Duplicate:    ATVACIDtdu      atvacid[e]      <=>      atvacid[c]
   Keep:    OXYPthc        oxyp[e]        <=>      oxyp[c]
Duplicate:    OXYPtepv      oxyp[c]        <=>      oxyp[e]
   Keep:    PVSHtu        pvs[e]        <=>      pvs[c]
Duplicate:    PVStep        pvs[c]        <=>      pvs[e]

```

Remove any duplicate reactions from the stoichiometric matrix, but do not remove the protons.

```

if length(removedRxnInd)>0
  irrevFlag=0;
  metFlag=0;%dont remove the protons
  model = removeRxns(model,model.rxns(removedRxnInd),irrevFlag,metFlag);
end

```

Display statistics of the removed reactions

```

if printLevel>0
  [nMet0,nRxn0]=size(modelOrig.S);
  [nMet,nRxn]=size(model.S);
  fprintf( '%6s\t%6s\n', '#mets', '#rxns' )
  fprintf( '%6u\t%6u\t%s\n', nMet0,nRxn0, ' totals.' )
  fprintf( '%6u\t%6u\t%s\n', nMet0-nMet,nRxn0-nRxn, ' duplicate reactions
upto protons removed.' )
  fprintf( '%6u\t%6u\t%s\n', nMet,nRxn, ' remaining.' )
end

```

```

#mets      #rxns
8399      13543      totals.
0          253       duplicate reactions upto protons removed.
8399      13290      remaining.

```

```

%model size
[nMet,nRxn]=size(model.S);

```

Heuristically identify exchange reactions and metabolites exclusively involved in exchange reactions

An external reaction is one that is heuristically identified by a single stoichiometric coefficient in the corresponding column of S, or an (abbreviated) reaction name matching a pattern (e.g. prefix EX_) or an external subsystem assignment. Any remaining reaction is assumed to be an internal reaction. If a reaction is not external then it is denoted an internal reaction. External reactants are exclusively involved in exchange reactions, and internal reactants otherwise. The findSExRxnInd function finds the external reactions in the model which export or import mass from or to the model, e.g. Exchange reactions, Demand reactions, Sink reactions.

```

if ~isfield(model,'SIntMetBool') || ~isfield(model,'SIntRxnBool')
  model = findSExRxnInd(model,[],printLevel-1);
end

```

Assuming biomass reaction is: biomass_reaction

EXPECTED RESULTS

In the returned model, `model.SIntRxnBool`, is a boolean of reactions heuristically thought to be mass balanced, while `model.SIntMetBool` is a boolean of metabolites heuristically thought to be involved in mass balanced reactions.

CAUTION

The aforementioned assignments of external and internal reactions and reactants is the result of a heuristic and might result in one or more errors, either due to misspecification or because the names of external reactions and external subsystems often vary between laboratories.

Find the reactions that are flux inconsistent

Ultimately we seek to identify the set of stoichiometrically consistent reactions that are also flux consistent, with no bounds on reaction rates. However, finding the stoichiometrically consistent subset can be demanding for large models so first we identify the subset of reactions that are flux consistent and focus on them.

```
modelOrig=model;
model.lb(~model.SIntRxnBool)=-1000;
model.ub(~model.SIntRxnBool)= 1000;
if 1
    if ~isfield(model,'fluxConsistentMetBool') || ~isfield(model,'fluxConsistentRxnBool')
        param.modeFlag=0;
        param.method='null_fastcc';
        %param.method='fastcc';
        [fluxConsistentMetBool,fluxConsistentRxnBool,...]
            fluxInConsistentMetBool,fluxInConsistentRxnBool,model]...
            = findFluxConsistentSubset(model,param,printLevel);
    end
    % Remove reactions that are flux inconsistent
    if any(fluxInConsistentRxnBool)
        irrevFlag=0;
        metFlag=1;
        model =
removeRxns(model,model.rxns(fluxInConsistentRxnBool),irrevFlag,metFlag);
        [nMet0,nRxn0]=size(modelOrig.S);
        [nMet,nRxn]=size(model.S);

        if printLevel>0
            fprintf('%s\n','-----')
            fprintf('%6s\t%6s\n','#mets','#rxns')
            fprintf('%6u\t%6u\t%$s\n',nMet0,nRxn0,' totals.')
            fprintf('%6u\t%6u\t%$s\n',nMet0-nMet,nRxn0-nRxn,' flux
inconsistent reactions removed.')
            fprintf('%6u\t%6u\t%$s\n',nMet,nRxn,' remaining.')
            fprintf('%s\n','-----')
```

```

        if printLevel>1
            for n=1:nRxn0
                if fluxInConsistentRxnBool(n)

fprintf('%.15s\t%-100s\n',modelOrig.rxns{n},modelOrig.rxnNames{n})
            end
        end
    end
%revise model size
[nMet,nRxn]=size(model.S);

%Recompute
%Heuristically identify exchange reactions and metabolites
exclusively involved in exchange reactions
%finds the reactions in the model which export/import from the model
%boundary i.e. mass unbalanced reactions
%e.g. Exchange reactions
%     Demand reactions
%     Sink reactions

model = findSExRxnInd(model,[],0);
if printLevel>0
    fprintf('%s\n','-----end-----')
end
end
end

```

```

--- findFluxConsistentSubset START ----
12164      Total reactions
5974       Reversible reactions.
6190       Irreversible reactions.
6777 flux consistent metabolites
1622 flux inconsistent metabolites
11802 flux consistent reactions
1488 flux inconsistent reactions
--- findFluxConsistentSubset END ----
-----
#mets      #rxns
8399      13290      totals.
1622       1488       flux inconsistent reactions removed.
6777      11802      remaining.
-----
3HPCOAHYD   3-Hydroxyisobutyryl-Coenzyme A Hydrolase
3HPPD       3-Hydroxypropionate Dehydrogenase
3NTD71      3'-Nucleotidase (AMP), Lysosomal
4MPTNLtr    4-Methylpentanal Transport, Endoplasmatic Reticulum
5HOXINDACTOXm 5-Hydroxyindoleacetaldehyde:NAD+ Oxidoreductase, Mitochondrial
A_MANASE    Alpha-Mannosidase
ACSOMT     S-Adenosyl-L-Methionine:N-Acetylserotonin O-Methyltransferase
ADEt1       Adenine Faciliated Transport from Lysosome
ADPGLC     ADPglucose Diphosphatase
ADPRDPm    ADPribose Diphosphatase, Mitochondrial
ADSELK     Adenylyl-Selenate Kinase
AGLPR      Alkyl Glycerol Phosphate Reductase
AGPex      Alkyl Glycerol Phosphate Transport

```

AGPRim	N-Acetyl-G-Glutamyl-Phosphate Reductase, Irreversible, Mitochondrial
AGPSx	Alkylglycerone Phosphate Synthase
ALKP	Alkaline Phosphatase
ALOX12R	Arachidonate 12-Lipoxygenase R
AMACR2r	Alpha-Methylacyl Coenzyme A Racemase (Reductase)
AMACRr	Alpha-Methylacyl Coenzyme A Racemase
AMPtr	AMP Transporter, Endoplasmic Reticulum
AP4AH1	Ap4A Hydrolase, Asymmetrically
BAMPALDOXm	Beta-Aminopropion Aldehyde:NAD+ Oxidoreductase, Mitochondrial
BDG2HCGHD	Beta-D-Glucosyl-2-Coumarinate Glucohydrolase
C2M26DCO AHLm	Cis-2-Methyl-5-Isopropylhexa-2, 5-Dienoyl Coenzyme A Hydro-Lyase, Mitochondrial
C2M26DCO AHLx	Cis-2-Methyl-5-Isopropylhexa-2, 5-Dienoyl Coenzyme A Hydro-Lyase, Peroxisomal
CBR1	Carbonyl Reductase [NADPH] 1
CCA_D3t	Calcitroic Acid Transport from Cytosol
CCA_D3tm	Calcitroic Acid Transport from Mitochondria
CO2tn	CO2 Nuclear Transport via Diffusion
CPCTDTX	Choline-Phosphate Cytidylyltransferase
CRTSTRNtr	Corticosterone Intracellular Transport
CYSLYSL	L-Cystine Lysteine-Lyase (Deaminating)
CYSTAm	Cysteine Transaminase, Mitochondrial
DALAt2rL	D-Alanine Transport via Proton Symport, Lysosomal
DEDOLP1_U	Dehydrodolichol Diphosphate Phosphatase (Uterus)
DEDOLP2_U	Dehydrodolichol Phosphate Phosphatase (Uterus)
DEDOLR_U	Dehydrodolichol Reductase (Uterus)
DHAPAx	Dihydroxyacetone Phosphate Acyltransferase
DMHPTCRNCPT1	Carnitine Fatty-Acyl Transferase
DOGULND1	2, 3-Dioxo-L-Gulonate Decarboxylase (L-Lyxonate-Forming)
DOGULND2	2, 3-Dioxo-L-Gulonate Decarboxylase (L-Xylonate-Forming)
DOGULNO2	2, 3-Dioxo-L-Gulonate:Hydrogen Peroxide Oxireductase
DPROOp	D-Proline Oxidase, Peroxisomal
ECGISOr	Ecgongine Isomerase, Endoplasmatic Reticulum
EGMESTr	Ecgongine Methyl Esterase, Endoplasmatic Reticulum
ENGASE	Endo-Beta-N-Acetylglucosaminidase
ENGASE2	Endo-Beta-N-Acetylglucosaminidase
ENMAN1g	Endomannosidase (Glc1Man-Producing), Golgi Apparatus
ENMAN2g	Endomannosidase (Glc2Man-Producing), Golgi Apparatus
ENMAN3g	Endomannosidase (Glc3Man-Producing), Golgi Apparatus
ENMAN4g	Endomannosidase (M6Masnc-Producing), Golgi Apparatus
ENMAN5g	Endomannosidase (M6Masnb2-Producing), Golgi Apparatus
ENMAN6g	Endomannosidase (M5Masnb1-Producing), Golgi Apparatus
EPCTX	Ethanolamine-Phosphate Cytidylyltransferase
EX_cca_d3[e]	Exchange of Calcitroic Acid (D3)
EX_pro_D[e]	Exchange of D-Proline
EX_sel[e]	Exchange of Selenate
EX_ser_D[e]	Exchange of D-Serine
EX_vitd2[e]	Exchange of Vitamin D2
FA120ACPH	Fatty-Acyl-Acp Hydrolase
FA140ACPH	Fatty-Acyl-Acp Hydrolase
FA141ACPH	Fatty-Acyl-Acp Hydrolase
FA161ACPH	Fatty-Acyl-Acp Hydrolase
FA180ACPH	Fatty-Acyl-Acp Hydrolase
FA181ACPH	Fatty-Acyl-Acp Hydrolase
FA1821ACPH	Fatty-Acyl-Acp Hydrolase
FA1822ACPH	Fatty-Acyl-Acp Hydrolase
FA182ACPH	Fatty-Acyl-Acp Hydrolase
G1M6MASNB1terg	Transport of Glucosyl-(Alpha-D-Mannosyl)6-Beta-D-Mannosyl-Diacetylchitobiosyl-L-Asparag
G1M7MASNBterg	Transport of Glucosyl-(Alpha-D-Mannosyl)7-Beta-D-Mannosyl-Diacetylchitobiosyl-L-Asparag
G1M7MASNCterg	Transport of Glucosyl-(Alpha-D-Mannosyl)7-Beta-D-Mannosyl-Diacetylchitobiosyl-L-Asparag
G1M8MASNterg	Transport of (Alpha-D-Glucosyl)-(Alpha-D-Mannosyl)8-Beta-D-Mannosyl-Diacetylchitobiosyl
G2M8MASNterg	Transport of (Alpha-D-Glucosyl)2-(Alpha-D-Mannosyl)8-Beta-D-Mannosyl-Diacetylchitobiosyl
G3M8MASNterg	Transport of (Alpha-D-Glucosyl)3-(Alpha-D-Mannosyl)8-Beta-D-Mannosyl-Diacetylchitobiosyl
GGT_U	Geranylgeranyltransferase (Uterus)
GHMT3m	Glycine Hydroxymethyltransferase, Mitochondrial
GK1m	Guanylate Kinase (GMP:ATP), Mitochondrial

GLACOm	D-Glucuronolactone:NAD+ Oxidoreductase, Mitochondrial
GPAMm_hs	Glycerol-3-Phosphate Acyltransferase
GSNKm	Guanosine Kinase, Mitochondrial
GSNtm	Guanosine Faciliated Transport in Mitochondria
H8MTer_L	H8 Mannosyltransferase, Endoplasmic Reticulum
H8MTer_U	H8 Mannosyltransferase, Endoplasmic Reticulum
HEXCCPT2	Carnitine Transferase
HEXCCRNT	Transport into the Mitochondria (Carnitine)
HMGCOArr	Hydroxymethylglutaryl Coenzyme A Reductase (Ir)
HXANTl	Hypoxanthine Faciliated Transport from Lysosome
IMACTD_m	Imidazole Acetaldeyde Dehydrogenase, Mitochondrial
INSKm	Insosine Kinase, Mitochondrial
INSTl	Transport of Inosine, Faciliated, Lysosomal
INStm	Transport of Inosine, Faciliated, Mitochondrial
IPDPtr	Isopentenyl Diphosphate Transport, Endoplasmatic Reticulum
LACZly	B-Galactosidase, Lysosomal
LCADI_Dm	Lactaldehyde Dehydrogenase, Mitochondrial
LCTStl	Lactose Transport from Cytosol to Lysosome (Via Autophagocytosis)
LCYSTATm	L-Cysteate:2-Oxoglutarate Aminotransferase, Mitochondrial
LGNCCPT2	Transport into the Mitochondria (Carnitine)
LGNCCRNT	Transport into the Mitochondria (Carnitine)
LS3	Lumisterol 3 Formation
LYSMTF1n	Histone-Lysine N-Methyltransferase, Nuclear
LYSMTF2n	Histone-Lysine N-Methyltransferase, Nuclear
LYSMTF3n	Histone-Lysine N-Methyltransferase, Nuclear
M4ATAer	M4A Transamidase, Endoplasmic Reticulum
M4BET2er	M4B Phosphoethanolaminyl Transferase, Endoplasmic Reticulum
MAN1_6B1er	Mannosidase I, Endoplasmic Reticulum (G1M6Masnb1-Producing)
MAN1_7Ber	Mannosidase I, Endoplasmic Reticulum (G1M7Masnb-Producing)
MAN2_6B1er	Mannosidase II, Endoplasmic Reticulum (G1M6Masnb1-Producing)
MAN2_7Cer	Mannosidase II, Endoplasmic Reticulum (G1M7Masnc-Producing)
MCOATAm	Malonyl Coenzyme A-Acp Transacylase, Mitochondrial
MEOHtr	Methanol Transporter, Endoplasmic Reticulum
MI1345PKn	Inositol-1, 3, 4, 5-Triphosphate 6-Kinase, Nucleus
MI1346PKn	Inositol-1, 3, 4, 6-Tetrakisphosphate 5-Kinase, Nucleus
MI1346Ptn	1D-Myo-Inositol 1, 3, 4, 6-Tetrakisphosphate Nuclear Transport (Diffusion)
MI134PK	Inositol-1, 3, 4-Trisphosphate 6-Kinase
MI1456PKn	Inositol-1, 4, 5, 6- Tetrakisphosphate 3-Kinase, Nucleus
MI145P6Kn	Inositol-1, 4, 5-Triphosphate 6-Kinase, Nucleus
MI145PKn	Inositol-1, 4, 5-Trisphosphate 3-Kinase, Nucleus
MI3456PK	Inositol-3, 4, 5, 6-Tetrakisphosphate 1-Kinase
MMCD	Methylmalonyl Coenzyme A Decarboxylase
MMCDp	Methylmalonyl Coenzyme A Decarboxylase, Peroxisomal
NABTNOm	N4-Acetylaminobutanal:NAD+ Oxidoreductase, Mitochondrial
NDPK10m	Nucleoside-Diphosphate Kinase (ATP:DIDP), Mitochondrial
NDPK2m	Nucleoside-Diphosphate Kinase (ATP:UDP), Mitochondrial
NDPK9m	Nucleoside-Diphosphate Kinase (ATP:IDP), Mitochondrial
NMPTRCOX	N-Methylputrescine:Oxygen Oxidoreductase (Deaminating)
NNATm	Nicotinate-Nucleotide Adenylyltransferase, Mitochondrial
NRVNCCPT2	Carnitine Transferase
NRVNCCRNT	Transport into the Mitochondria (Carnitine)
NTD2m	5'-Nucleotidase (UMP), Mitochondrial
NTD31	5'-Nucleotidase (dCMP), Lysosomal
NTD61	5'-Nucleotidase (dAMP), Lysosomal
NTD81	5'-Nucleotidase (dGMP), Lysosomal
NTMELYStner	Protein Trimethyl Lysine Transport (Nucleus to Endoplasmatic Reticulum)
NTPP11	Nucleoside Triphosphate Pyrophosphorylase (XTP)
PE_HStg	Phosphatidylethanolamine Scramblase
PECGONCOATr	Pseudoecgonine Coenzyme A Transferase, Endoplasmatic Reticulum
PEPLYStn	Peptide (Lysine) Nuclear Transport via Diffusion
PI45PLCn	Phosphatidylinositol 4, 5-Bisphosphate Phospholipase C, Nucleus
PI4P3Ker	Phosphatidylinositol 4-Phosphate 3-Kinase, Endoplasmic Reticulum
PI5P3Ker	Phosphatidylinositol-5-Phosphate 3-Kinase, Endoplasmic Reticulum
PIK3er	Phosphatidylinositol 3-Kinase, Endoplasmic Reticulum

PLYSPSer	Protein Lysine Peptidase (Endoplasmic Reticulum)
PNTKm	Pantothenate Kinase, Mitochondrial
PROAKGOX1r	L-Proline, 2-Oxoglutarate:Oxygen Oxidoreductase (4-Hydroxylating), Endoplasmatic Reticulum
PRODt2r	D-Proline Reversible Transport via Proton Symport
PRODt2rL	D-Proline Reversible Transport via Proton Symport, Lysosomal
PRPNCOAHYDx	Propenoyl Coenzyme A Hydrolase, Peroxisomal
PS_HStg	Phosphatidylserine Scramblase
PTE5x	Peroxisomal Acyl Coenzyme A Thioesterase
PYAM5Ptm	Pyridoxamine 5'-Phosphate Transport via Diffusion, Mitochondrial
PYDX5Ptm	Pyridoxal 5'-Phosphate Transport via Diffusion, Mitochondrial
PYLALDOX	Perillyl Aldehyde:NAD+ Oxidoreductase
PYLALDOXm	Perillyl Aldehyde:NAD+ Oxidoreductase, Mitochondrial
RETNCOA	Retinoyl Coenzyme A Formation
Rtotaltp	Fatty Acid Intracellular Transport
SELADT	Selenate Adenylyltransferase
SELCYSLY	Selenocysteine Lyase
SELCYSLY2	Selenocysteine Lyase
SELNPS	Selenophosphate Synthase
SELt4_3	Selenate Transport via Sodium Symport
SGPL11r	Sphingosine-1-Phosphate Lyase 1
SIAASE	Sialidase
SLDxm	L-Sulfolactate Dehydrogenase (NAD+), Mitochondrial
SOAT11r	Sterol O-Acyltransferase (Acyl-Coenzyme A: Cholesterol Acyltransferase) 1
SOAT12r	Sterol O-Acyltransferase (Acyl-Coenzyme A: Cholesterol Acyltransferase) 1
SRTN23OX	5-Hydroxytryptamine:Oxygen 2, 3-Dioxygenase (Indole-Decyclizing)
SRTNMTX	S-Adenosyl-L-Methionine:Amine N-Methyltransferase (Srtm)
T2M26DCO AHLm	Trans-2-Methyl-5-Isopropylhexa-2, 5-Dienoyl Coenzyme A Hydro-Lyase, Mitochondrial
T2M26DCO AHLx	Trans-2-Methyl-5-Isopropylhexa-2, 5-Dienoyl Coenzyme A Hydro-Lyase, Peroxisomal
T4HCINNOX	4-Coumarate:Oxygen Oxidoreductase
TDPDRR	DTDP-4-Dehydrorhamnose Reductase
TMLYSter	Trimethyl-L-Lysine Transport (Er to Cytosol)
TRDRm	Thioredoxin Reductase (NADPH)
UDPGALT2g	UDPGalactose Transport, Golgi Apparatus
UDPGLCtg	UDP-Glc Golgi Transport via CMP Antiport
UGALNACTer	UDP-GalacNAc Endoplasmic Reticulum Transport via CMP Antiport
UMPKm	UMP Kinase (Mitochondrial, ATP)
Uritm	Uridine Facilitated Transport in Mitochondria
VITD2Hm	Vitamin D-25-Hydroxylase (D2)
VITD2t	Vitamin D2 Release
VITD2tm	Vitamin D2 Transport from Mitochondria
XOL7AH2tr	Lipid, Flip-Flop Intracellular Transport
XOLDIOLONETm	Lipid, Flip-Flop Intracellular Transport
r0001	Virtual Reaction/Potential Definition
r0120	GTP 7, 8-8, 9-Dihydrolase
r0121	GTP 7, 8-8, 9-Dihydrolase
r0267	CMP-N-Acetylneuraminate, Ferrocyanochrome-B5:Oxygen Oxidoreductase (N-Acetyl-Hydroxylating)
r0268	Cytidine Monophospho-N-Acetylneurameric Acid Hydroxylase
r0400	N-Acetylneuraminate, Ferrocyanochrome-B5:Oxygen Oxidoreductase (N-Acetyl-Hydroxylating)
r0598	L-Fucose Ketol-Isomerase
r0625	3Alpha, 7Alpha, 12Alpha-Trihydroxy-5Beta-Cholestan-26-Al: NAD+ 26-Oxidoreductase Bile Acid
r0626	5Beta-Cholestane-3Alpha, 7Alpha, 12Alpha, 26-Tetraol: NAD+ 26-Oxidoreductase Bile Acid
r0668	CTP:N-Acetylneuraminate Cytidyllyltransferase
r0678	Acyl-[Acyl-Carrier-Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
r0681	(3R)-3-Hydroxybutanoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis
r0682	Butyryl-[Acyl-Carrier Protein]:Malonyl Coenzyme A C-Acyltransferase (Decarboxylating, C)
r0691	(3R)-3-Hydroxybutanoyl-[Acyl-Carrier Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
r0692	(3R)-3-Hydroxydecanoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
r0693	(3R)-3-Hydroxybutanoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis
r0694	(3R)-3-Hydroxyoctanoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
r0695	(3R)-3-Hydroxybutanoyl-[Acyl-Carrier-Protein] Hydro-Lyase
r0696	(3R)-3-Hydroxypalmitoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
r0697	(3R)-3-Hydroxypalmitoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis
r0701	(3R)-3-Hydroxytetradecanoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Bi
r0702	(3R)-3-Hydroxypalmitoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis

r0708 2-Amino-4-Hydroxy-6- (Erythro-1, 2, 3-Trihydroxypropyl) Dihydropteridine Triphosphate 7
 r0709 2-Amino-4-Hydroxy-6- (Erythro-1, 2, 3-Trihydroxypropyl) Dihydropteridine Triphosphate 7
 r0712 Dodecanoyl-[Acyl-Carrier Protein]: Malonyl Coenzyme A C-Acyltransferase (Decarboxylating)
 r0713 Dodecanoyl-[Acyl-Carrier-Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0760 Butyryl-[Acyl-Carrier Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0761 (3R)-3-Hydroxyhexanoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
 r0762 (3R)-3-Hydroxybutanoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis
 r0763 Hexanoyl-[Acyl-Carrier Protein]:Oxoacyl- And Enoyl-Reducing And Thioester-Hydrolysing
 r0764 Hexanoyl-[Acyl-Carrier Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0765 Octanoyl-[Acyl-Carrier Protein]:Malonyl Coenzyme A C-Acyltransferase (Decarboxylating,
 r0766 Octanoyl-[Acyl-Carrier Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0767 Decanoyl-[Acyl-Carrier Protein]:Malonyl Coenzyme A C-Acyltransferase (Decarboxylating,
 r0768 Decanoyl-[Acyl-Carrier Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0769 (3R)-3-Hydroxylodecanoyl-[Acyl-Carrier-Protein]:NADP+ Oxidoreductase Fatty Acid Biosynthesis
 r0770 (3R)-3-Hydroxybutanoyl-[Acyl-Carrier-Protein] Hydro-Lyase Fatty Acid Biosynthesis
 r0771 Tetradecanoyl-[Acyl-Carrier Protein]:Malonyl Coenzyme A C-Acyltransferase (Decarboxylating)
 r0772 Tetradecanoyl-[Acyl-Carrier Protein]:Malonyl-[Acyl-Carrier-Protein] C-Acyltransferase (Decarboxylating)
 r0773 Hexadecanoyl-[Acyl-Carrier Protein]:Malonyl Coenzyme A C-Acyltransferase (Decarboxylating)
 r0775 Formamidopyrimidine Nucleoside Triphosphate 7, 8-8, 9-Dihydrolase
 r0776 Formamidopyrimidine Nucleoside Triphosphate 7, 8-8, 9-Dihydrolase
 r0777 GTP 7, 8-8, 9-Dihydrolase
 r0778 GTP 7, 8-8, 9-Dihydrolase
 r0800 Virtual ReactionPotential Definition
 r0988 Postulated Transport Reaction
 r0992 Na (+)Bile Acid Symporter Active Transport
 r1021 Postulated Transport Reaction
 r1027 Active Transport
 r1131 Transport Reaction
 r1132 Transport Reaction
 r1133 Transport Reaction
 r1319 Virtual ReactionPotential Definition
 r1320 Virtual ReactionPotential Definition
 r1321 Virtual ReactionPotential Definition
 r1322 Virtual ReactionPotential Definition
 r1323 Virtual ReactionPotential Definition
 r1324 Virtual ReactionPotential Definition
 r1325 Virtual ReactionPotential Definition
 r1326 Virtual ReactionPotential Definition
 r1327 Virtual ReactionPotential Definition
 r1328 Virtual ReactionPotential Definition
 r1329 Virtual ReactionPotential Definition
 r1330 Virtual ReactionPotential Definition
 r1331 Virtual ReactionPotential Definition
 r1332 Virtual ReactionPotential Definition
 r1430 [Acyl-Carrier-Protein] 4-Pantetheine-Phosphohydrolase
 r1431 2-Deoxyuridine 5-Diphosphate:Oxidized-Thioredoxin 2-Oxidoreductase
 r1432 2-Deoxyuridine 5-Diphosphate:Oxidized-Thioredoxin 2-Oxidoreductase
 r1441 Active Transport
 r1526 ATP-Binding Cassette (ABC) Tcdb:3.A.1.211.1
 r2073 Zinc (Zn²⁺)-Iron (Fe²⁺) Permease (Zip), Tcdb:2.A.55.2.3
 RE0066C Phosphatidylethanolamine N-Methyltransferase
 RE0066M Phosphatidylethanolamine N-Methyltransferase
 RE0066R Phosphatidylethanolamine N-Methyltransferase
 RE0344M Palmitoyl Coenzyme A Hydrolase
 RE0344X Palmitoyl Coenzyme A Hydrolase
 RE0452N DTMP Kinase
 RE0456M Ribonucleoside-Diphosphate Reductase
 RE0512C 3-Hydroxyacyl Coenzyme A Dehydrogenase
 RE0572N RE0572N
 RE0573N RE0573N
 RE0577M Palmitoyl Coenzyme A Hydrolase
 RE0577X Palmitoyl Coenzyme A Hydrolase
 RE0578M Palmitoyl Coenzyme A Hydrolase
 RE0578X Palmitoyl Coenzyme A Hydrolase

RE0579M	Palmitoyl Coenzyme A Hydrolase
RE0579X	Palmitoyl Coenzyme A Hydrolase
RE0580L	RE0580L
RE0580R	RE0580R
RE0581R	RE0581R
RE0582N	RE0582N
RE0583N	RE0583N
RE0688C	RE0688C
RE0688X	RE0688X
RE0689C	RE0689C
RE0689X	RE0689X
RE0690C	RE0690C
RE0690X	RE0690X
RE0702C	Dihydrolipoyl Dehydrogenase
RE0702L	Dihydrolipoyl Dehydrogenase
RE0702M	Dihydrolipoyl Dehydrogenase
RE0702N	Dihydrolipoyl Dehydrogenase
RE0827C	RE0827C
RE0827X	RE0827X
RE0828C	RE0828C
RE0828X	RE0828X
RE0864C	RE0864C
RE0875C	RE0875C
RE0908G	Steryl-Sulfatase
RE0908R	Steryl-Sulfatase
RE0916G	Steryl-Sulfatase
RE0916R	Steryl-Sulfatase
RE0918G	Steryl-Sulfatase
RE0918R	Steryl-Sulfatase
RE0919C	Glucuronosyltransferase
RE0919R	Glucuronosyltransferase
RE0920C	Glucuronosyltransferase
RE0920R	Glucuronosyltransferase
RE0921C	Glucuronosyltransferase
RE0921R	Glucuronosyltransferase
RE0922C	Glucuronosyltransferase
RE0922R	Glucuronosyltransferase
RE0923C	Glucuronosyltransferase
RE0923R	Glucuronosyltransferase
RE0924C	Glucuronosyltransferase
RE0924R	Glucuronosyltransferase
RE0925C	Glucuronosyltransferase
RE0925R	Glucuronosyltransferase
RE0926C	Alpha-Amylase
RE0927C	Glucuronosyltransferase
RE0927R	Glucuronosyltransferase
RE0928C	Glucuronosyltransferase
RE0928R	Glucuronosyltransferase
RE0935C	Alpha-Amylase
RE0936C	RE0936C
RE0937C	RE0937C
RE0938C	RE0938C
RE0944C	Alpha-Amylase
RE1050C	Peroxidase
RE1050L	Peroxidase
RE1050N	Peroxidase
RE1062C	Neurolysin
RE1062M	Neurolysin
RE1063C	Thimet Oligopeptidase
RE1064C	Thimet Oligopeptidase
RE1096M	RE1096M
RE1096R	RE1096R
RE1099G	Steryl-Sulfatase
RE1099L	Steryl-Sulfatase

RE1099R	Steryl-Sulfatase
RE1100G	Steryl-Sulfatase
RE1100L	Steryl-Sulfatase
RE1134M	RE1134M
RE1134R	RE1134R
RE1135G	Steryl-Sulfatase
RE1135L	Steryl-Sulfatase
RE1233M	Kynurenine-Oxoglutarate Transaminase
RE1236C	RE1236C
RE1238X	Diamine N-Acetyltransferase
RE1240C	RE1240C
RE1317C	L-Iditol 2-Dehydrogenase
RE1441G	1-Phosphatidylinositol-4-Phosphate 5-Kinase
RE1473C	Gamma-Glutamyltransferase
RE1508C	RE1508C
RE1514M	Long-Chain-Fatty-Acid- Coenzyme A Ligase
RE1514X	Long-Chain-Fatty-Acid- Coenzyme A Ligase
RE1525C	3-Hydroxyacyl Coenzyme A Dehydrogenase
RE1526C	3-Hydroxyacyl Coenzyme A Dehydrogenase
RE1527C	3-Hydroxyacyl Coenzyme A Dehydrogenase
RE1537C	RE1537C
RE1537X	RE1537X
RE1538C	RE1538C
RE1538X	RE1538X
RE1539C	RE1539C
RE1539X	RE1539X
RE1582L	Quinine 3-Monooxygenase
RE1587L	Quinine 3-Monooxygenase
RE1651C	NADPH:Quinone Reductase
RE1653C	NADPH:Quinone Reductase
RE1711M	Alcohol Dehydrogenase
RE1796C	Steroid Delta-Isomerase
RE1806C	Quinine 3-Monooxygenase
RE1809C	Quinine 3-Monooxygenase
RE1809R	Quinine 3-Monooxygenase
RE1811C	Quinine 3-Monooxygenase
RE1811R	Quinine 3-Monooxygenase
RE1812C	Quinine 3-Monooxygenase
RE1812R	Quinine 3-Monooxygenase
RE1818C	Glutathione Transferase
RE1818M	Glutathione Transferase
RE1818R	Glutathione Transferase
RE1818X	Glutathione Transferase
RE1819C	Carbonyl Reductase (NADPH)
RE1819M	Carbonyl Reductase (NADPH)
RE1819X	Carbonyl Reductase (NADPH)
RE1826M	RE1826M
RE1827M	RE1827M
RE1828C	RE1828C
RE1828M	RE1828M
RE1829C	RE1829C
RE1829M	RE1829M
RE1830C	RE1830C
RE1830M	RE1830M
RE1835M	Palmitoyl Coenzyme A Hydrolase
RE1835X	Palmitoyl Coenzyme A Hydrolase
RE1836M	Propionyl Coenzyme A C2-Trimethyltridecanoyltransferase
RE1846X	Bile Acid-CoA:Amino Acid N-Acyltransferase
RE1860C	2',3'-Cyclic-Nucleotide 3'-Phosphodiesterase
RE1899C	Deoxyhypusine Synthase
RE1907C	RE1907C
RE1916X	Glutathione Transferase
RE1917C	RE1917C
RE1920C	Catechol O-Methyltransferase

RE1922C	Catechol O-Methyltransferase
RE1927C	RE1927C
RE1942C	RE1942C
RE1942R	RE1942R
RE1952C	Microsomal Epoxide Hydrolase
RE1952R	Microsomal Epoxide Hydrolase
RE1952X	Microsomal Epoxide Hydrolase
RE1954C	RE1954C
RE1956C	Microsomal Epoxide Hydrolase
RE1956R	Microsomal Epoxide Hydrolase
RE1956X	Microsomal Epoxide Hydrolase
RE1957G	1-Phosphatidylinositol-4-Phosphate 5-Kinase
RE2026C	RE2026C
RE2027C	RE2027C
RE2028C	RE2028C
RE2029C	RE2029C
RE2031M	Amino-Acid N-Acetyltransferase
RE2032M	Amino-Acid N-Acetyltransferase
RE2040C	Gamma-Glutamylcyclotransferase
RE2041C	Gamma-Glutamylcyclotransferase
RE2048N	Arachidonate 5-Lipoxygenase
RE2048R	Arachidonate 5-Lipoxygenase
RE2049C	RE2049C
RE2050C	Prostaglandin-Endoperoxide Synthase
RE2050R	Prostaglandin-Endoperoxide Synthase
RE2051C	Phosphatidate Phosphatase
RE2051G	Phosphatidate Phosphatase
RE2051R	Phosphatidate Phosphatase
RE2067C	RE2067C
RE2068C	RE2068C
RE2070C	RE2070C
RE2081C	Peptide Alpha-N-Acetyltransferase
RE2117M	Glycine N-Acyltransferase
RE2124C	Catechol O-Methyltransferase
RE2128C	RE2128C
RE2129C	RE2129C
RE2131C	RE2131C
RE2133C	Catechol O-Methyltransferase
RE2138C	RE2138C
RE2139C	RE2139C
RE2140C	RE2140C
RE2141C	RE2141C
RE2146C	Glucuronosyltransferase
RE2146R	Glucuronosyltransferase
RE2149C	Glucuronosyltransferase
RE2149R	Glucuronosyltransferase
RE2150C	Glucuronosyltransferase
RE2150R	Glucuronosyltransferase
RE2152C	RE2152C
RE2155R	Steroid 21-Monoxygenase
RE2156M	Amino-Acid N-Acetyltransferase
RE2202C	RE2202C
RE2203C	RE2203C
RE2221C	RE2221C
RE2221M	RE2221M
RE2248C	RE2248C
RE2250C	RE2250C
RE2251C	RE2251C
RE2252C	RE2252C
RE2265C	Tissue Kallikrein
RE2269C	Chymase
RE2270C	Carboxypeptidase A
RE2272C	Tripeptidyl-Peptidase I
RE2272L	Tripeptidyl-Peptidase I

```

RE2273C Carboxypeptidase A
RE2292C RE2292C
RE2296X Glutathione Transferase
RE2306C Pyroglutamyl-Peptidase II
RE2333C RE2333C
RE2334C RE2334C
RE2335C RE2335C
RE2349M Kynurenine-Oxoglutarate Transaminase
RE2360C RE2360C
RE2360N RE2360N
RE2373C RE2373C
RE2375C RE2375C
RE2377C RE2377C
RE2384C RE2384C
RE2404R Glucuronosyltransferase
RE2405R Glucuronosyltransferase
RE2410N 7-Dehydrocholesterol Reductase
RE2440C RE2440C
RE2443C Glutathione Transferase
RE2443M Glutathione Transferase
RE2444C RE2444C
RE2445C Peptidyl-Dipeptidase A
RE2452C RE2452C
RE2453M Methylcrotonoyl Coenzyme A Carboxylase
RE2454M Methylcrotonoyl Coenzyme A Carboxylase
RE2459C Sterol Esterase
RE2474C Quinine 3-Monooxygenase
RE2474R Quinine 3-Monooxygenase
RE2476C RE2476C
RE2477C RE2477C
RE2493C Methionine Synthase
RE2513C Peroxidase
RE2513L Peroxidase
RE2513N Peroxidase
RE2514C Peroxidase
RE2514L Peroxidase
RE2514N Peroxidase
RE2520C NADPH:Quinone Reductase
RE2521C NADPH:Quinone Reductase
RE2522C Glutathione Transferase
RE2522X Glutathione Transferase
RE2523C Glutathione Transferase
RE2523X Glutathione Transferase
RE2524C Glutathione Transferase
RE2524X Glutathione...

```

Found biomass reaction: biomass_reaction

-----end-----

Find mass leaks or siphons within the heuristically internal part, without using the bounds given by the model

```

if 1
    modelBoundsFlag=0;
    leakParams.method='dc';
    leakParams.theta=0.5;

[leakMetBool,leakRxnBool,siphonMetBool,siphonRxnBool,leaky,siphony,statp,statn] = ...
    findMassLeaksAndSiphons(model,model.SIntMetBool,model.SIntRxnBool,...)

```

```

modelBoundsFlag, leakParams, printLevel);
end

6776    9958      subset tested for leakage (dc method, with infinite flux bounds)...
5028    6528      semipositive leaking metabolites (and exclusive reactions).
983.289 10fthf5glu[c]
983.289 10fthf5glu[l]
983.289 10fthf5glu[m]
1111.32  10fthf6glu[c]
1111.32  10fthf6glu[l]
1111.32  10fthf6glu[m]
1239.36  10fthf7glu[c]
1239.36  10fthf7glu[l]
1239.36  10fthf7glu[m]
471.15   10fthf[c]
...
11DOCRTSLtm
11DOCRTSLtr
11DOCRTSTRNtm
11DOCRTSTRNtr
13DAMPPOX
1MNCAMti
1PPDCRp
24_25DHVITD3t
24_25DHVITD3tm
25HVITD3t
  5028    6528      seminegative siphon metabolites (and exclusive reactions).
983.289 10fthf5glu[c]
983.289 10fthf5glu[l]
983.289 10fthf5glu[m]
1111.32  10fthf6glu[c]
1111.32  10fthf6glu[l]
1111.32  10fthf6glu[m]
1239.36  10fthf7glu[c]
1239.36  10fthf7glu[l]
1239.36  10fthf7glu[m]
471.15   10fthf[c]
...
11DOCRTSLtm
11DOCRTSLtr
11DOCRTSTRNtm
11DOCRTSTRNtr
13DAMPPOX
1MNCAMti
1PPDCRp
24_25DHVITD3t
24_25DHVITD3tm
25HVITD3t

```

Find the maximal set of reactions that are stoichiometrically consistent

```

if ~isfield(model, 'SConsistentMetBool') ||
~isfield(model, 'SConsistentRxnBool')
    if strcmp(model.modelID, 'HMRdatabase2_00')
        massBalanceCheck=0;
    else
        massBalanceCheck=1;
    end
    if 1

```

```

[SConsistentMetBool,SConsistentRxnBool,SInConsistentMetBool,SInConsistentRxnBool,
ool,unknownSConsistencyMetBool,unknownSConsistencyRxnBool,model]...
    =findStoichConsistentSubset(model,massBalanceCheck,printLevel);
else
    %print out problematic reactions to file
    resultsFileName=[resultsPath filesep model.modelID];

[SConsistentMetBool,SConsistentRxnBool,SInConsistentMetBool,SInConsistentRxnBool,
ool,unknownSConsistencyMetBool,unknownSConsistencyRxnBool,model]...

=findStoichConsistentSubset(model,massBalanceCheck,printLevel,resultsFileName
);
end
end

```

```

--- findStoichConsistentSubset START ----
-----
#mets      #rxns
6777      11802      totals.

-----
1          1844      heuristically external.
6776      9958      heuristically internal.
3676      9238      seemingly elementally balanced.
3676      9238      seemingly elementally balanced and stoichiometrically consistent.
3101      2564      seemingly elementally imbalanced.

-----
3676      9238      heuristically internal and seemingly elementally balanced.
3676      9238      seemingly elementally balanced and stoichiometrically consistent.
3100      720       heuristically internal and seemingly elementally imbalanced.

-----
Iteration #1 minimum cardinality of conservation relaxation vector.
6776      9958      unknown consistency.
6776      9958      being tested.
6776      9475      ... of which are stoichiometrically consistent by min cardinality of stoich consistency.

Infeasibility while detecting semipositive leaking metabolites.
6776      9475      Confirmed stoichiometrically consistent by leak/siphon testing.
0          483       ... of which are of unknown consistency.
0          101       removed.

-----
Iteration #2 minimum cardinality of conservation relaxation vector.
0          382       unknown consistency.
6776      9857      being tested.
6776      9601      ... of which are stoichiometrically consistent by min cardinality of stoich consistency.
6776      9601      Confirmed stoichiometrically consistent by leak/siphon testing.
0          256       ... of which are of unknown consistency.
0          22        removed.

-----
Iteration #3 minimum cardinality of conservation relaxation vector.
0          234       unknown consistency.
6776      9835      being tested.
6776      9605      ... of which are stoichiometrically consistent by min cardinality of stoich consistency.

Infeasibility while detecting seminegative leaking metabolites.
6776      9605      Confirmed stoichiometrically consistent by leak/siphon testing.
0          230       ... of which are of unknown consistency.
0          8         removed.

-----
Iteration #4 minimum cardinality of conservation relaxation vector.
0          222       unknown consistency.
6776      9827      being tested.

```

```

6776      9610      ... of which are stoichiometrically consistent by min cardinality of stoich consistent
Infeasibility while detecting semipositive leaking metabolites.
6776      9610      Confirmed stoichiometrically consistent by leak/siphon testing.
0          217      ... of which are of unknown consistency.
0          3          removed.
-----
Iteration #5 minimum cardinality of conservation relaxation vector.
0          214      unknown consistency.
6776      9824      being tested.
6776      9609      ... of which are stoichiometrically consistent by min cardinality of stoich consistent
Infeasibility while detecting seminegative leaking metabolites.
6776      9609      Confirmed stoichiometrically consistent by leak/siphon testing.
0          215      ... of which are of unknown consistency.
-----
Iteration #6 minimum cardinality of conservation relaxation vector.
0          215      unknown consistency.
6776      9824      being tested.
6776      9609      ... of which are stoichiometrically consistent by min cardinality of stoich consistent
Infeasibility while detecting seminegative leaking metabolites.
6776      9609      Confirmed stoichiometrically consistent by leak/siphon testing.
0          215      ... of which are of unknown consistency.
Infeasibility while detecting seminegative leaking metabolites.
--- Summary of stoichiometric consistency ---
6777      11802      totals.
1          1844      heuristically external.
6776      9958      heuristically internal:
6776      9609      ... of which are stoichiometrically consistent.
0          134      ... of which are stoichiometrically inconsistent.
0          215      ... of which are of unknown consistency.
---
0          349      heuristically internal and stoichiometrically inconsistent or unknown consistency.
644        134      ... of which are elementally imbalanced (inclusively involved metabolite).
0          134      ... of which are elementally imbalanced (exclusively involved metabolite).
6776      9609      Confirmed stoichiometrically consistent by leak/siphon testing.
--- findStoichConsistentSubset END ---

```

```

rxnBool=model.SInConsistentRxnBool & model.SIntRxnBool;
if any(rxnBool)
    if printLevel>0
        fprintf('%s\n','Stoichiometrically inconsistent heuristically non-
exchange reactions:')
    end
    for n=1:nRxn
        if rxnBool(n)
            fprintf('%20s\t%50s\t%s\n',model.rxns{n},model.rxnNames{n})
        end
    end
    if printLevel>0
        fprintf('%s\n','-----')
    end
end

```

Stoichiometrically inconsistent heuristically non-exchange reactions:

ABO7g Abo Blood Group (Transferase A, Alpha 1-3-N-Acetylgalactosaminyltransferase

```

rxnBool=model.unknownSConsistencyRxnBool & model.SIntRxnBool;
if any(rxnBool)
    if printLevel>0
        fprintf('%s\n', 'Unknown consistency heuristically non-exchange
reactions:')
    end
    for n=1:nRxn
        if rxnBool(n)
            fprintf('%20s\t%50s\t%s\n',model.rxns{n},model.rxnNames{n})
        end
    end
    if printLevel>0
        fprintf('%s\n', '-----')
    end
end

```

```

Unknown consistency heuristically non-exchange reactions:
    3MOBt2im      3-Methyl-2-Oxobutanoate Mitochondrial Transport via Proton Symport
-----

```

Sanity check of stoichiometric and flux consistency of model with open external reactions

```

if all(model.SIntMetBool & model.SConsistentMetBool)...
    && nnz(model.SIntRxnBool &
model.SConsistentRxnBool)==nnz(model.SIntRxnBool)...
    && all(model.fluxConsistentMetBool)...
    && all(model.fluxConsistentRxnBool)

[nMet,nRxn]=size(model.S);
if printLevel>1
    fprintf('%6s\t%6s\n', '#mets', '#rxns')
    fprintf('%6u\t%6u\t%s\n', nMet, nRxn, ' totals.')

fprintf('%6u\t%6u\t%s\n', nnz(~model.SIntMetBool), nnz(~model.SIntRxnBool), 'heuristically exchange.')
end

checksPassed=0;
%Check that all heuristically non-exchange reactions are also
stoichiometrically consistent

%exchange reactions
model.EXRxnBool=strncmp('EX_', model.rxns, 3)==1;
%demand reactions going out of model
model.DMRxnBool=strncmp('DM_', model.rxns, 3)==1;
%sink reactions going into or out of model
model.SinkRxnBool=strncmp('sink_', model.rxns, 5)==1;
%all heuristic non-exchanges, i.e., supposedly all external reactions

```

```

        bool=~(model.EXRxnBool | model.DMRxnBool | model.SinkRxnBool);
        if nnz(bool & model.SIntRxnBool &
model.SConsistentRxnBool)==nnz(model.SConsistentRxnBool)
            checksPassed=checksPassed+1;
            if printLevel>1

fprintf('%6u\t%6u\t%s\n',nnz(model.SIntMetBool),nnz(model.SIntRxnBool),' All
internally stoichiometrically consistent. (Check 1: minimum cardinality of
conservation relaxation vector.)');
            end
        end

        %Check for mass leaks or siphons in the stoichiometrically
consistent part
        %There should be no leaks or siphons in the stoichiometrically
consistent part
        modelBoundsFlag=0;
        leakParams.epsilon=1e-4;
        leakParams.eta = getCobraSolverParams('LP', 'feasTol')*100;
        leakParams.method='dc';

[leakMetBool,leakRxnBool,siphonMetBool,siphonRxnBool,leaky,siphony,statp,stat
n]...

=findMassLeaksAndSiphons(model,model.SConsistentMetBool,model.SConsistentRxnB
ool,modelBoundsFlag,leakParams,printLevel);

        if nnz(leakMetBool)==0 && nnz(leakRxnBool)==0 &&
nnz(siphonMetBool)==0 && nnz(siphonRxnBool)==0
            checksPassed=checksPassed+1;
            if printLevel>1
                fprintf('%6u\t%6u\t%s\n',nnz(leakMetBool |
siphonMetBool),nnz(leakRxnBool | siphonRxnBool),' No internal leaks or
siphons. (Check 2: leak/siphon tests.');
                end
            end

        %Check that the maximal conservation vector is nonzero for each the
%internal stoichiometric matrix
        maxCardinalityConsParams.epsilon=1e-4;%1/epsilon is the largest mass
considered, needed for numerical stability
        maxCardinalityConsParams.method = 'quasiConcave';%seems to work the
best, but sometimes infeasible
        maxCardinalityConsParams.theta = 0.5;
        maxCardinalityConsParams.eta=getCobraSolverParams('LP',
'feasTol')*100;

[maxConservationMetBool,maxConservationRxnBool,solution]=maxCardinalityConser
vationVector(model.S(model.SConsistentMetBool,model.SConsistentRxnBool),
maxCardinalityConsParams);

```

```

    if nnz(maxConservationMetBool)==size(model.S,1) &&
nnz(maxConservationRxnBool)==nnz(model.SIntRxnBool)
        checksPassed=checksPassed+1;
        if printLevel>1

fprintf('%6u\t%6u\t%s\n',nnz(maxConservationMetBool),nnz(maxConservationRxnBo
ol),' All internally stoichiometrically consistent. (Check 3: maximim
cardinality conservation vector.)');
        end
    end

    %Check that each of the reactions in the model (with open external
reactions) is flux consistent
    modelOpen=model;
    modelOpen.lb(~model.SIntRxnBool)=-1000;
    modelOpen.ub(~model.SIntRxnBool)= 1000;
    param.epsilon=1e-4;
    param.modeFlag=0;
    param.method='null_fastcc';

[fluxConsistentMetBool,fluxConsistentRxnBool,fluxInConsistentMetBool,fluxInCo
nsistentRxnBool,modelOpen] =
findFluxConsistentSubset(modelOpen,param,printLevel-2);

    if nnz(fluxConsistentMetBool)==size(model.S,1) &&
nnz(fluxConsistentRxnBool)==size(model.S,2)
        checksPassed=checksPassed+1;
        if printLevel>1

fprintf('%6u\t%6u\t%s\n',nnz(fluxConsistentMetBool),nnz(fluxConsistentRxnBool
),' All flux consistent. (Check 4: maximim cardinality constrained right
nullspace.)');
        end
    end

    if checksPassed==4
        %save the model with open exchanges as the default generic
        %model
        model=modelOpen;
        if printLevel>0
            fprintf('%s\n','Open external reactions is
stoichiometrically and flux consistent. A flux balance model generated from
a reconstruction. GREAT!!!!');
            end
        end
        save([resultsFileName '_consistent.mat'],'model')
    end

```

REFERENCES

- Gevorgyan, A., Poolman, M. G., Fell D., Detection of stoichiometric inconsistencies in biomolecular models. *Bioinformatics*, 24(19):2245–51, 2008.
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