

Screen Records

Now that the Exclusions Reasons have been configured, you can proceed with screening underlying studies to identify those that should be Included for your nest, or Excluded (for one of your configured Exclusion Reasons).

Note: If you are using Two-Pass Screening or Dual Screening, this process will differ slightly from the Standard workflow outlined below. See the [Two-Pass Screening](#), [Dual Screening](#), [Dual Two-Pass Screening](#) pages for more details!

Steps for Standard Screening:

1. Navigate to Screening

You can either Screen Sequentially (by selecting “Screening” in the menu, outlined in red below), where records will be shown to you in order of expected Inclusion Probability, or screen from Inspector (outlined in black).

The screenshot displays the Nested Knowledge web application interface. On the left is a sidebar menu with the following items: 'Nest Home', 'Activity Settings', 'Literature Search' (with sub-items 'Other Sources', 'Duplicate Review', 'Search Exploration'), 'Screening' (highlighted with a red border and a gear icon), 'Tagging' (with a gear icon), 'Study Inspector', and 'Synthesis' (with sub-items 'Manuscript Editor', 'Abstract Editor', 'Export'). The main content area is titled 'Protocol' and shows a 'COVID-19: Antivirals (Demo)' protocol. It includes an 'About' section, a 'Title' section, a 'Study Coordinator/Corresponding Author' section with contact information for Erin Sheffels, and a 'Team Members and Their Organizational Affiliations' section listing various researchers. On the right is a 'Notes' panel showing a conversation between Kathryn Cowie and Karl Holub.

2. Read study abstract

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Suemori, 2021

Abstract

Full Text

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Related Reports

PMC

A multicenter non-randomized, uncontrolled single arm trial for evaluation of the efficacy and the safety of the treatment with favipiravir for patients with severe fever with thrombocytopenia syndrome.

Severe fever with thrombocytopenia syndrome (SFTS) is a bunyavirus infection with high mortality. Favipiravir has shown effectiveness in preventing and treating SFTS virus (SFTSV) infection in animal models. A multicenter non-randomized, uncontrolled single arm trial was conducted to collect data on the safety and the effectiveness of favipiravir in treatment of SFTS patients. All participants received favipiravir orally (first-day loading dose of 1800 mg twice a day followed by 800 mg twice a day for 7-14 days in total). SFTSV RT-PCR and biochemistry tests were performed at designated time points. Outcomes were 28-day mortality, clinical improvement, viral load evolution, and adverse events (AEs). Twenty-six patients were enrolled, of whom 23 were analyzed. Four of these 23 patients died of multi-organ failure within one week (28-day mortality rate: 17.3%). Oral favipiravir was well tolerated in the surviving patients. AEs (abnormal hepatic function and insomnia) occurred in about 20% of the patients. Clinical symptoms improved in all patients who survived from a median of day 2 to day10. SFTSV RNA levels in the patients who died were significantly higher than those in the survivors (p = 0.0029). No viral genomes were detectable in the surviving patients a median of 8 days after favipiravir administration. The 28-day mortality rate in this study was lower than those of the previous studies in Japan. The high frequency of hepatic dysfunction as an AE was observed. However, it was unclear whether this was merely a side effect of favipiravir, because liver disorders are commonly seen in SFTS patients. The results of this trial support the effectiveness of favipiravir for patients with SFTS.

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Screening

Full Text Review

P(Inclusion): 0.77

Upload Full Text

Exclude:

Search Reasons

Select Reason

Not an RCT of a drug of interest

Protocol or Methods article

Systematic Review or Meta-analysis

Editorial, comment, or opinion article

Not related to COVID-19

Update or guidelines article

Qualitative review of existing research

Include:

Include

Tagging

Comments (0)

History

Your task in screening should be to identify, based on the Abstract content, whether the record falls under any Exclusion Reason, or whether it is on-topic for your review and satisfies your criteria for inclusion.

The Screening page displays an abstract highlighted withRoboPICO, which is an open source fork of the models offered in RobotReviewer that identifies the Population, Interventions, and Outcomes in an abstract. Then, see on the right a panel to select Exclusion Reasons or Include the article in question.

Using the scite banner

Above your abstract, you can see the scite banner, which displays the number of times the publication in question was cited, supported, mentioned, and contrasted. If you click the banner, you can see more citation-related information provided by scite.ai, including retractions!

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scite_

Smart Citations

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Citing Publications

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Supporting

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Contrasting

View Citations

See how this article has been cited at scite.ai

scite shows how a scientific paper has been cited by providing the context of the citation, a classification describing whether it supports, mentions, or contrasts the cited claim, and a label indicating in which section the citation was made.

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3. Decide if study should be Included or Excluded

If the abstract does not provide enough information for you to decide if it should be Included or Excluded, click on the study source button (in this case PubMed, see red arrow below) and source the full text of the study.

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If you read the FULL TEXT and decide it should be included, check the “Full Text Review” box.

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Exclude Records

If you read the abstract and find that one or more of your Exclusion Reasons (red box above) are applicable, click on the reason that applies to that specific study. This will apply your reason and automatically bring up the next study to be screened.

Include Records

If you read the abstract and find that none of your Exclusion Reasons apply, and that (based on information available to you) the publication in question is relevant to your review, select “Include” (see red box above).

Skipping a study

Having a hard time deciding whether to include or exclude a study? You can hit skip and leave it unscreened until you're ready to make a decision.

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Add Exclusion Reasons on the Fly

You can add Exclusions Reasons as you screen without leaving the Screening page. To do so, in the Screening module, open the Exclusion Reason drop-down and begin typing in an Exclusion Reason.

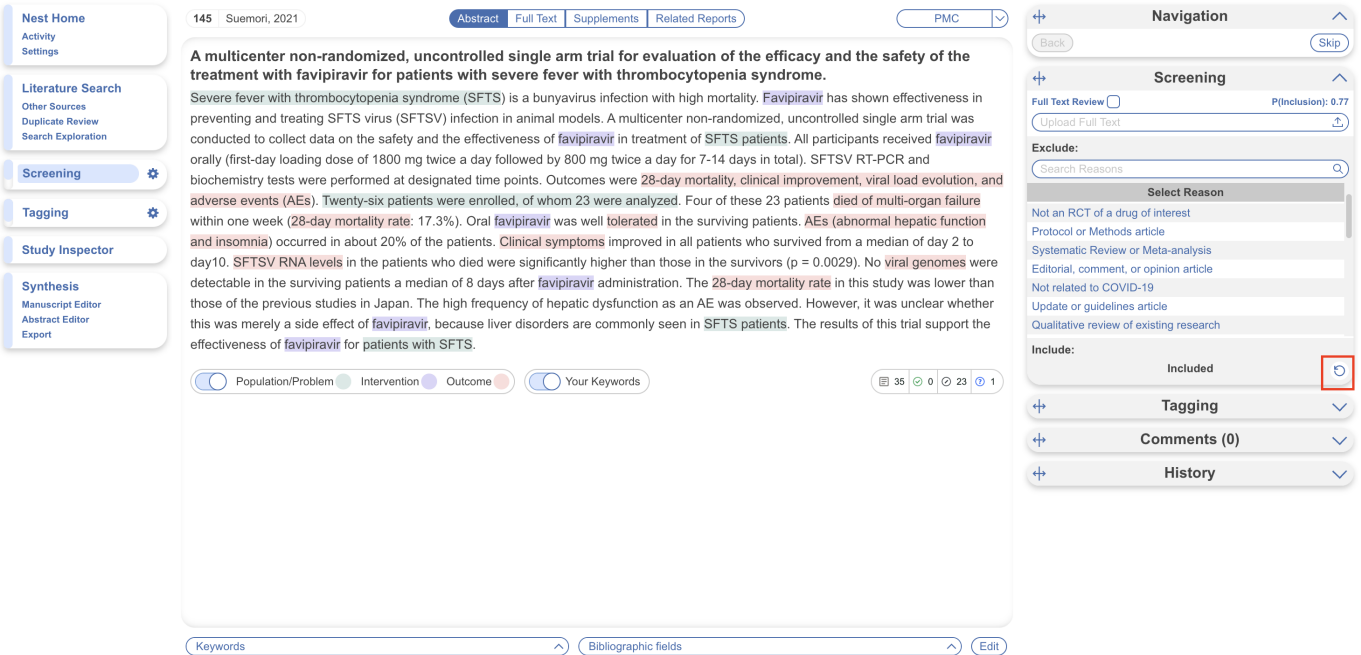
If the reason of interest has not yet been configured, you will be presented with the ability to “Add Option.” Select this option, and write out your full Exclusion Reason. Once you have added it, it will be added to the Exclusion Reason drop-down and the Configure Exclusion Reasons page, and will be automatically applied to the study you are currently screening. To confirm that the new reason should

be applied, select “Exclude”.

Unscreening a study

If you have included or excluded a study that you want to revert to 'unscreened' status so that it can be reviewed again, you can unscreen it by finding the study of interest in [Study Inspector](#), and then selecting the icon next to the Include button on the study you want to unscreen. A pop-up will appear and you can then click “Unscreen” to unscreen that single study.

Note: if you want to unscreen multiple studies, you can also do so using [Bulk Actions](#)!



Note: Anytime there is a module box with the adjustable icon, you can drag to adjust the width of the box depending on your preference.

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Included

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Comments (0)

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4. Upload the Full Text

In general, uploading a Full Text should be completed only for Included records, and doing so assists in preparing the Tagging step.

For instructions on how to upload a Full Text PDF, click [here](#).

No Full Text

If you cannot source a full text for the study in question, you can use the “No Full Text” option to designate an Exclusion Reason specifically to address those records.

For those records, first configure an Exclusion Reason as “No Full Text” in the Configure Exclusion Reasons page:

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Add

Exclusion Reasons

Import Set

Reason		Excluded Records	No Full Text	
pediatrics		3	Signals No FT <input type="checkbox"/>	
Not Published in English		2	Signals No FT <input type="checkbox"/>	
Valsartan Heart Failure Trial		2	Signals No FT <input type="checkbox"/>	
Correspondence		1	Signals No FT <input type="checkbox"/>	
Based on retracted study		1	Signals No FT <input type="checkbox"/>	
ST-Segment Elevation Myocardial Infarction		1	Signals No FT <input type="checkbox"/>	
Reports patients with ejection fraction above 45 ...		1	Signals No FT <input type="checkbox"/>	
Not a pharmacological treatment		1	Signals No FT <input type="checkbox"/>	
No Ivabradine		1	Signals No FT <input type="checkbox"/>	
No full text		0	Signals No FT <input checked="" type="checkbox"/>	

Then, apply this Exclusion Reason to all records where a full text was sought but not found.

Implications: Marking “No Full Text” is a special PRISMA category, so the specific reason you configure for this purpose will be given its own listing in your [PRISMA chart](#).

5. Upload Supplementary Materials

If you want to upload supplementary files to a specific record, you can do so in the Supplements tab. To upload supplements, [follow these instructions](#).

6. Mark Related Reports

If you come across several studies as related to one another, you can mark it as a related report in the Related Reports tab. Then, the software will automatically adjust the PRISMA diagram to reflect this. To mark a paper as a related report, [follow these instructions](#).

7. Continue Screening

Once you have clicked “Include” or “Exclude” (or “skip”) for any study, you should be automatically shown the next study.

If you are screening from [Inspector](#), you can use the arrows in the far left and right of the screen to

navigate up or down, respectively, or click out to view the Inspector study list.

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