CONSORT-EHEALTH (V 1.6.1) -Submission/Publication Form

The CONSORT-EHEALTH checklist is intended for authors of randomized trials evaluating web-based and Internet-based applications/interventions, including mobile interventions, electronic games (incl multiplayer games), social media, certain telehealth applications, and other interactive and/or networked electronic applications. Some of the items (e.g. all subitems under item 5 - description of the intervention) may also be applicable for other study designs.

The goal of the CONSORT EHEALTH checklist and guideline is to be a) a guide for reporting for authors of RCTs, b) to form a basis for appraisal of an ehealth trial (in terms of validity)

CONSORT-EHEALTH items/subitems are MANDATORY reporting items for studies published in the Journal of Medical Internet Research and other journals / scientific societies endorsing the checklist.

Items numbered 1., 2., 3., 4a., 4b etc are original CONSORT or CONSORT-NPT (nonpharmacologic treatment) items. Items with Roman numerals (i., ii, iii, iv etc.) are CONSORT-EHEALTH extensions/clarifications.

As the CONSORT-EHEALTH checklist is still considered in a formative stage, we would ask that you also RATE ON A SCALE OF 1-5 how important/useful you feel each item is FOR THE PURPOSE OF THE CHECKLIST and reporting guideline (optional).

Mandatory reporting items are marked with a red *. In the textboxes, either copy & paste the relevant sections from your manuscript into this form - please include any quotes from your manuscript in QUOTATION MARKS,

or answer directly by providing additional information not in the manuscript, or elaborating on why the item was not relevant for this study.

YOUR ANSWERS WILL BE PUBLISHED AS A SUPPLEMENTARY FILE TO YOUR PUBLICATION IN JMIR AND ARE CONSIDERED PART OF YOUR PUBLICATION (IF ACCEPTED). Please fill in these questions diligently. Information will not be copyedited, so please use proper spelling and grammar, use correct capitalization, and avoid abbreviations.

DO NOT FORGET TO SAVE AS PDF _AND_ CLICK THE SUBMIT BUTTON SO YOUR ANSWERS ARE IN OUR DATABASE !!!

Citation Suggestion (if you append the pdf as Appendix we suggest to cite this paper in the caption):

Eysenbach G, CONSORT-EHEALTH Group CONSORT-EHEALTH: Improving and Standardizing Evaluation Reports of Web-based and Mobile Health Interventions J Med Internet Res 2011;13(4):e126 URL: <u>http://www.jmir.org/2011/4/e126/</u> doi: 10.2196/jmir.1923 PMID: 22209829

* Required

Your name * First Last

John Lalor

Primary Affiliation (short), City, Country * University of Toronto, Toronto, Canada

University of Notre Dame, Notre Dame, IN, USA

Your e-mail address * abc@gmail.com

john.lalor@nd.edu

Title of your manuscript * Provide the (draft) title of your manuscript.

Evaluating the Effectiveness of NoteAid in a Community Hospital Setting: Randomized Control Trial

Name of your App/Software/Intervention * If there is a short and a long/alternate name, write the short name first and add the long name in brackets.

ComprehENotes; NoteAid

Evaluated Version (if any) e.g. "V1", "Release 2017-03-01", "Version 2.0.27913"

Your answer

Language(s) *

What language is the intervention/app in? If multiple languages are available, separate by comma (e.g. "English, French")

English

URL of your Intervention Website or App

e.g. a direct link to the mobile app on app in appstore (itunes, Google Play), or URL of the website. If the intervention is a DVD or hardware, you can also link to an Amazon page.

https://bio-nlp.org/emrreadability/notesaid.uwm

URL of an image/screenshot (optional)	
Your answer	

Accessibility *

Can an enduser access the intervention presently?

- access is free and open
- access only for special usergroups, not open
- O access is open to everyone, but requires payment/subscription/in-app purchases
- o app/intervention no longer accessible
- Other:

Primary Medical Indication/Disease/Condition *

e.g. "Stress", "Diabetes", or define the target group in brackets after the condition, e.g. "Autism (Parents of children with)", "Alzheimers (Informal Caregivers of)"

Health Literacy

Primary Outcomes measured in trial * comma-separated list of primary outcomes reported in the trial

Improved EHR note comprehension

Secondary/other outcomes Are there any other outcomes the intervention is expected to affect?

Recommended "Dose" * What do the instructions for users say on how often the app should be used?
O Approximately Daily
O Approximately Weekly
O Approximately Monthly
O Approximately Yearly
"as needed"
O Other:

Approx. Percentage of Users (starters) still using the app as recommended after 3 months *

- unknown / not evaluated
- 0-10%
- 0 11-20%
- 0 21-30%
- 0 31-40%
- 0 41-50%
- 51-60%
- 61-70%
- 0 71%-80%
- 0 81-90%
- 91-100%
- O Other:

Overall, was the app/intervention effective? *
() yes: all primary outcomes were significantly better in intervention group vs control
$O _{\text{control}}^{\text{partly: SOME primary outcomes were significantly better in intervention group vs} $
O no statistically significant difference between control and intervention
$O\ $ potentially harmful: control was significantly better than intervention in one or more outcomes
O inconclusive: more research is needed
O Other:
Article Preparation Status/Stage * At which stage in your article preparation are you currently (at the time you fill in this form)

- not submitted yet in early draft status
- onot submitted yet in late draft status, just before submission
- Submitted to a journal but not reviewed yet
- Submitted to a journal and after receiving initial reviewer comments
- Submitted to a journal and accepted, but not published yet
- published
- O Other:

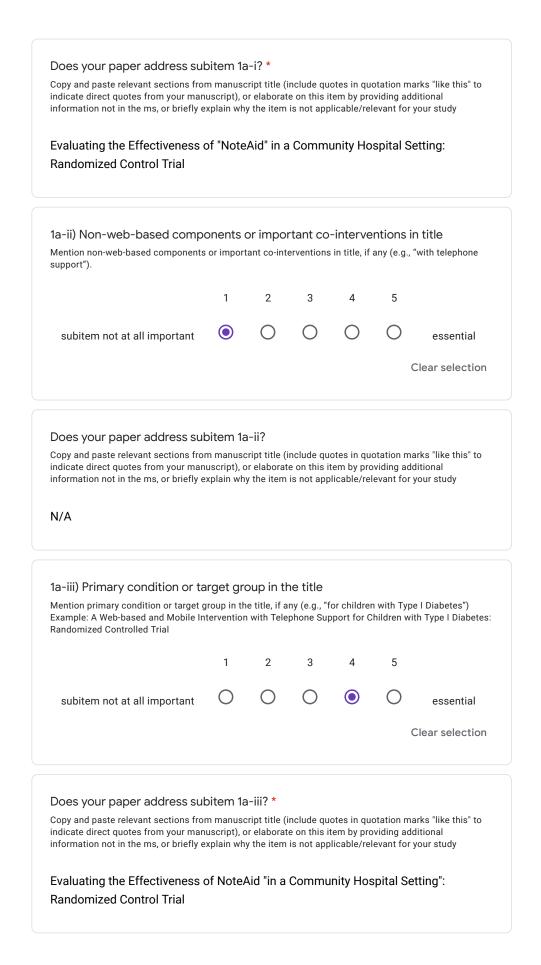
Journal *

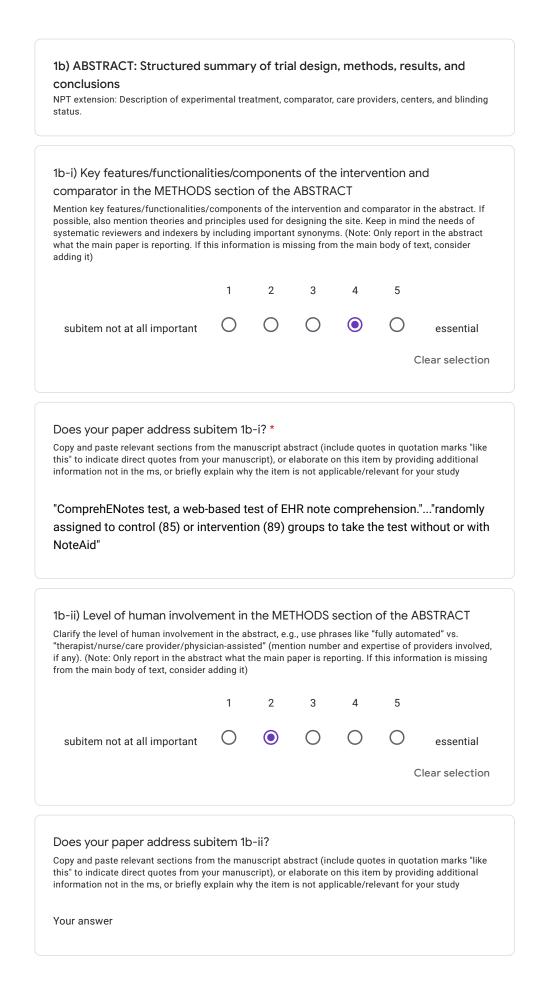
If you already know where you will submit this paper (or if it is already submitted), please provide the journal name (if it is not JMIR, provide the journal name under "other")

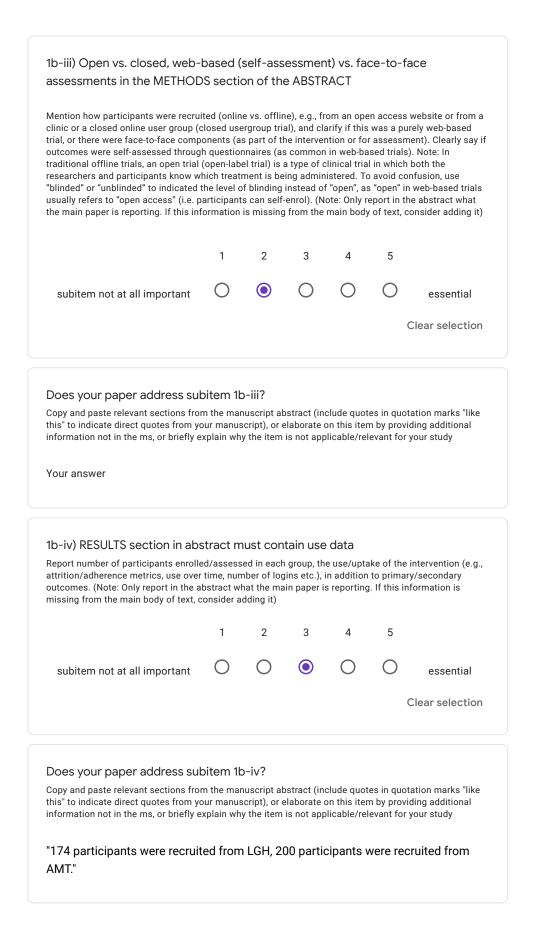
- not submitted yet / unclear where I will submit this
- Journal of Medical Internet Research (JMIR)
- JMIR mHealth and UHealth
- ◯ JMIR Serious Games
- JMIR Mental Health
- ◯ JMIR Public Health
- O JMIR Formative Research
- O ther JMIR sister journal
- O Other:

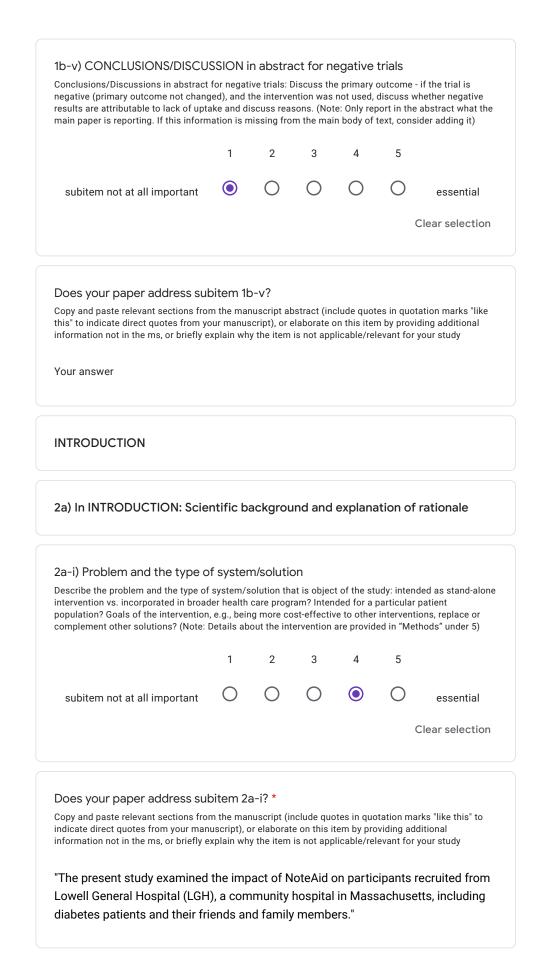
Is this a full powered effectiv	veness t	trial or a	pilot/fe	asibility	trial? *			
O Pilot/feasibility								
• Fully powered								
Manuscript tracking number	*							
If this is a JMIR submission, please provide the manuscript tracking number under "other" (The ms tracking number can be found in the submission acknowledgement email, or when you login as author in JMIR. If the paper is already published in JMIR, then the ms tracking number is the four-digit number at the end of the DOI, to be found at the bottom of each published article in JMIR)								
o ms number (yet) / not (y	et) subm	nitted to /	publishe	ed in JMI	R			
O Other:								
TITLE AND ABSTRACT								
1a) TITLE: Identification as a	randor	nized tr	ial in th	e title				
1a) Does your paper address I.e does the title contain the phrase " "other")				? (if not, ex	plain the re	eason under		
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I.e does the title contain the phrase " "other")	Randomiz	ed Control	led Trial"?	? (if not, ex	plain the re	eason under		
I.e does the title contain the phrase " "other") yes Other:	Randomiz livery ir bly use "w line", "virtu aponents li" only in groups". a as "mobi	the title veb-based ual", "intera (e.g. email the contex Compleme	ed Trial"?	nobile" and se "Interne mputer-bas al reality" (stitute prod	d/or "electi t-based" or sed" or "ele 3-D worlds duct name:	ronic game" in the nly if Intervention ectronic" only if e). Use "online" s with broader		
I.e does the title contain the phrase " "other") yes Other: 1a-i) Identify the mode of de Identify the mode of delivery. Prefera title. Avoid ambiguous terms like "on includes non-web-based Internet com offline products are used. Use "virtua only in the context of "online support terms for the class of products (such	Randomiz livery ir bly use "w line", "virtu aponents li" only in groups". a as "mobi	the title veb-based ual", "intera (e.g. email the contex Compleme	ed Trial"?	nobile" and se "Interne mputer-bas al reality" (stitute prod	d/or "electi t-based" or sed" or "ele 3-D worlds duct name:	ronic game" in the nly if Intervention ectronic" only if e). Use "online" s with broader		

Clear selection

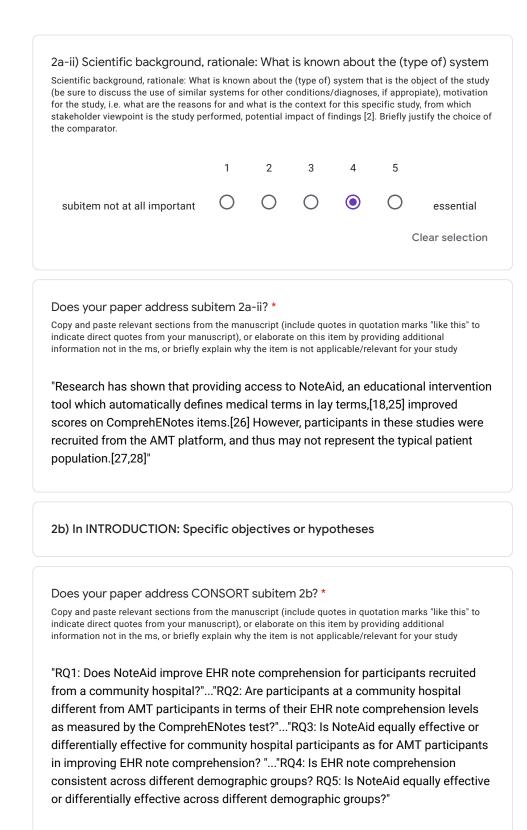






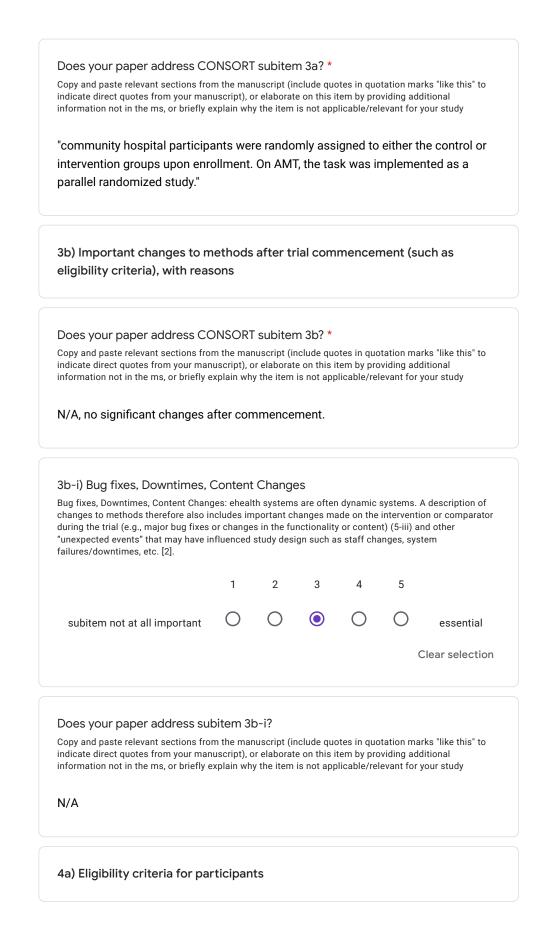


:



METHODS

3a) Description of trial design (such as parallel, factorial) including allocation ratio



Does your paper address CONSORT subitem 4a? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Patients, and persons accompanying patients in the waiting room, were eligible to participate if they were over 18, able to speak and read English, and comfortable using tablet."

 4a-i) Computer / Internet literacy

 Computer / Internet literacy is often an implicit "de facto" eligibility criterion - this should be explicitly clarified.

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 essential

Does your paper address subitem 4a-i?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Patients, and persons accompanying patients in the waiting room, were eligible to participate if they were over 18, able to speak and read English, and comfortable using tablet."

4a-ii) Open vs. closed, web-based vs. face-to-face assessments:

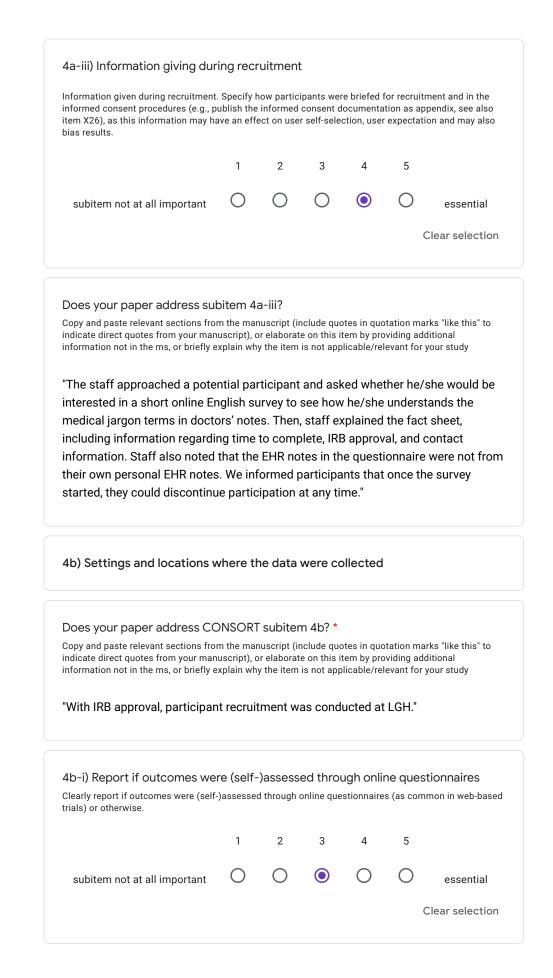
Open vs. closed, web-based vs. face-to-face assessments: Mention how participants were recruited (online vs. offline), e.g., from an open access website or from a clinic, and clarify if this was a purely webbased trial, or there were face-to-face components (as part of the intervention or for assessment), i.e., to what degree got the study team to know the participant. In online-only trials, clarify if participants were quasi-anonymous and whether having multiple identities was possible or whether technical or logistical measures (e.g., cookies, email confirmation, phone calls) were used to detect/prevent these.

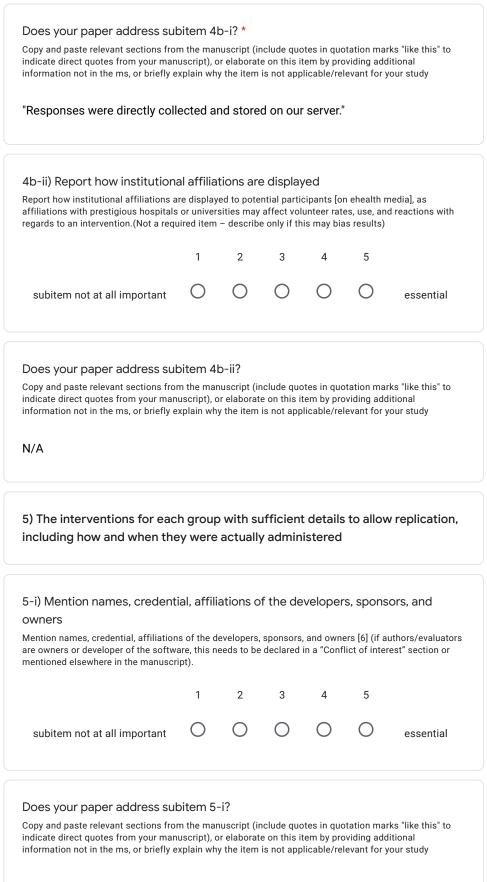
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Does your paper address subitem 4a-ii? *

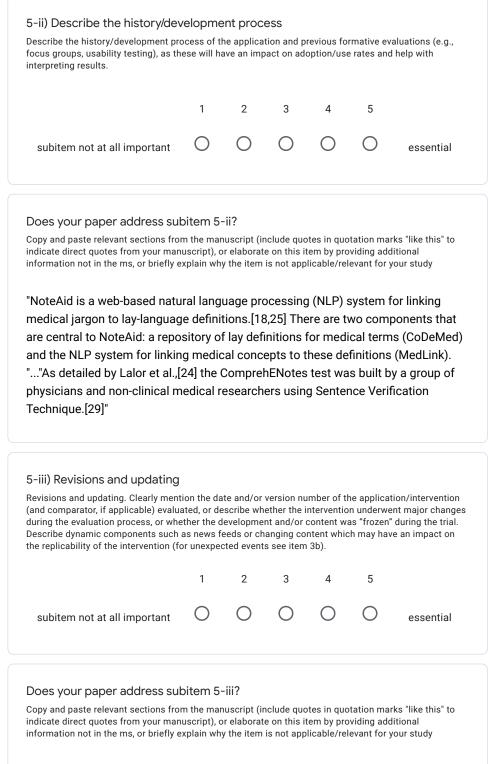
Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"With IRB approval, participant recruitment was conducted at LGH."

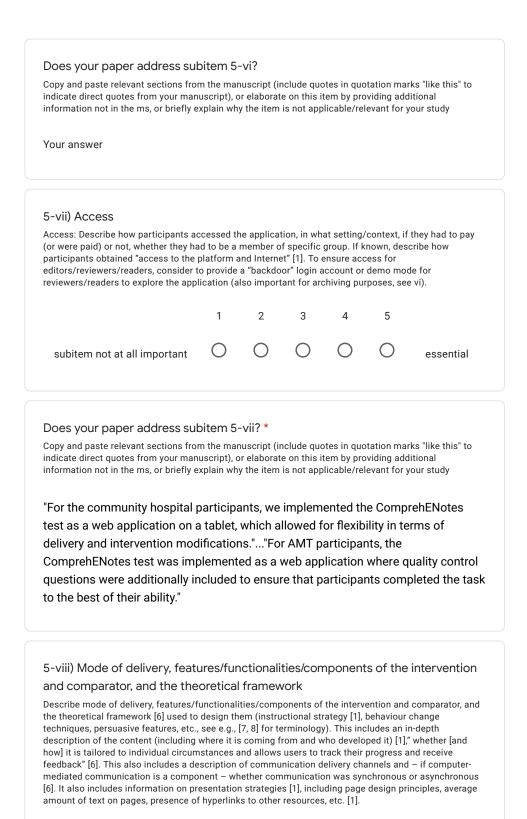




N/A



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5-v) Ensure replicability by post screenshots/screen-capture used Ensure replicability by publishing the and/or providing flowcharts of the alg principle be able to replicate the stud	video, a source co gorithms u	and/or p ode, and/or used. Repl	roviding providing icability (i.	g flowch I screensh e., other r	ots/screer	he algorithms		
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Does your paper address subitem 5-v? Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study Your answer								
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Does your paper address subitem 5-viii? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"For the intervention groups (community hospital and AMT), each ComprehENotes question was pre-processed by NoteAid and the results were embedded into the test web application directly. Definitions for terms were added to the web application as tooltip text. Terms were underlined to indicate that a definition was available, and when a participant hovered over a defined term the definition would automatically display (Figure 2). This behavior was also described in the introductory text paragraph of the web application so that the participants were aware of the definitions and knew how to access them."

5-ix) Describe use parameters

Describe use parameters (e.g., intended "doses" and optimal timing for use). Clarify what instructions or recommendations were given to the user, e.g., regarding timing, frequency, heaviness of use, if any, or was the intervention used ad libitum.

	1	2	3	4	5	
subitem not at all important	0	0	0	0	0	essential

Does your paper address subitem 5-ix?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

Your answer

5-x) Clarify the level of human involvement

Clarify the level of human involvement (care providers or health professionals, also technical assistance) in the e-intervention or as co-intervention (detail number and expertise of professionals involved, if any, as well as "type of assistance offered, the timing and frequency of the support, how it is initiated, and the medium by which the assistance is delivered". It may be necessary to distinguish between the level of human involvement required for the trial, and the level of human involvement required for a routine application outside of a RCT setting (discuss under item 21 – generalizability).

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subitem not at all important	0	0	0	0	0	essential

Does your paper address subitem 5-x?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

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subitem not at all important	0	0	0	0	0	essential
Does your paper address sul	oitem 5·	-xi? *				
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"For the intervention groups (commu	nity hos	pital and	d AMT),	each Co	mprehENote
question was pre-processed l	by Note	Aid and	the resu	lts were	embedo	led into the
test web application directly.						
application as tooltip text. Te						
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introductory text paragraph o aware of the definitions and k					e particit	ants were
5-xii) Describe any co-interv	entions	(incl. tra	aining/su	upport)		
5-xii) Describe any co-interv Describe any co-interventions (incl. tr addition to the targeted eHealth inter intervention. This includes training set the level of training required for the tr RCT setting (discuss under item 21 –	raining/su vention, a essions ar rial, and th	pport): Cle s ehealth i nd support ne level of	early state interventic [1]. It may	any interv on may not y be neces	be design sary to dis	ed as stand-alo tinguish betwee
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measures, including how and when they were assessed

Does your paper a	address CONSORT	subitem 6a? *
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Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"For our specific hypotheses regarding the effects of NoteAid in the two participant recruitment sources (hereafter referred to as "Source") (RQ1-3), we ran a two-way analysis of variance (ANOVA) to compare the four groups in our data set, using proportions of the passage-item pairs answered correctly as the dependent variable and Source (community hospital vs. AMT) and Condition (control vs. intervention) as two crossed factors."

6a-i) Online questionnaires: describe if they were validated for online use and apply CHERRIES items to describe how the questionnaires were designed/deployed

If outcomes were obtained through online questionnaires, describe if they were validated for online use and apply CHERRIES items to describe how the questionnaires were designed/deployed [9].

	1	2	3	4	5	
subitem not at all important	0	0	0	0	0	essential

Does your paper address subitem 6a-i? Copy and paste relevant sections from manuscript text

Your answer

6a-ii) Describe whether and how "use" (including intensity of use/dosage) was defined/measured/monitored

Describe whether and how "use" (including intensity of use/dosage) was defined/measured/monitored (logins, logfile analysis, etc.). Use/adoption metrics are important process outcomes that should be reported in any ehealth trial.

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subitem not at all important	0	0	0	0	0	essential

Does your paper address subitem 6a-ii?

Copy and paste relevant sections from manuscript text

6a-iii) Describe whether, was obtained Describe whether, how, and wher emails, feedback forms, interviev	n qualitative fe	eedback fro						
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Does your paper address Copy and paste relevant sections Your answer								
6b) Any changes to trial	outcomes	after th	ne trial c	ommen	ced, wi	th reasons		
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7a) How sample size was determined NPT: When applicable, details of whether and how the clustering by care provides or centers was addressed								
7a-i) Describe whether and how expected attrition was taken into account when calculating the sample size Describe whether and how expected attrition was taken into account when calculating the sample size.								
subitem not at all importar	1 nt O	2	3	4	5	essential		
Does your paper address subitem 7a-i? Copy and paste relevant sections from manuscript title (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study Your answer								

7b) When applicable, explanation of any interim analyses and stopping guidelines

Does your paper address CONSORT subitem 7b? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"We informed participants that once the survey started, they could discontinue participation at any time."

8a) Method used to generate the random allocation sequence NPT: When applicable, how care providers were allocated to each trial group

Does your paper address CONSORT subitem 8a? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Participants were randomly assigned to the control or treatment group when they accessed the web page to complete the test via a random number generator implemented in Python."

8b) Type of randomisation; details of any restriction (such as blocking and block size)

Does your paper address CONSORT subitem 8b? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"community hospital participants were randomly assigned to either the control or intervention groups upon enrollment. On AMT, the task was implemented as a parallel randomized study."

9) Mechanism used to implement the random allocation sequence (such as sequentially numbered containers), describing any steps taken to conceal the sequence until interventions were assigned

Does your paper address CONSORT subitem 9? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Participants were randomly assigned to the control or treatment group when they accessed the web page to complete the test via a random number generator implemented in Python."

10) Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions

Does your paper address CONSORT subitem 10? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Participants were randomly assigned to the control or treatment group when they accessed the web page to complete the test via a random number generator implemented in Python."

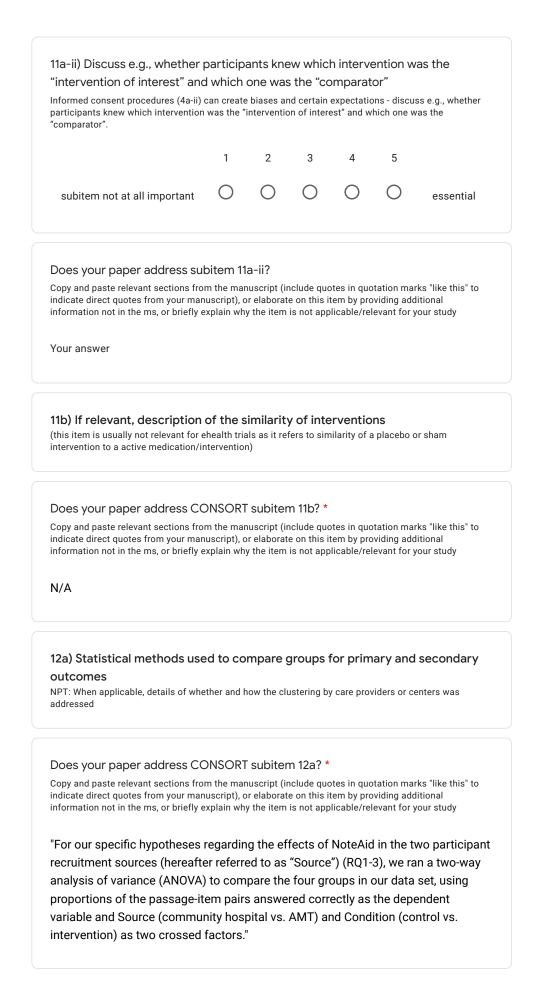
11a) If done, who was blinded after assignment to interventions (for example, participants, care providers, those assessing outcomes) and how NPT: Whether or not administering co-interventions were blinded to group assignment

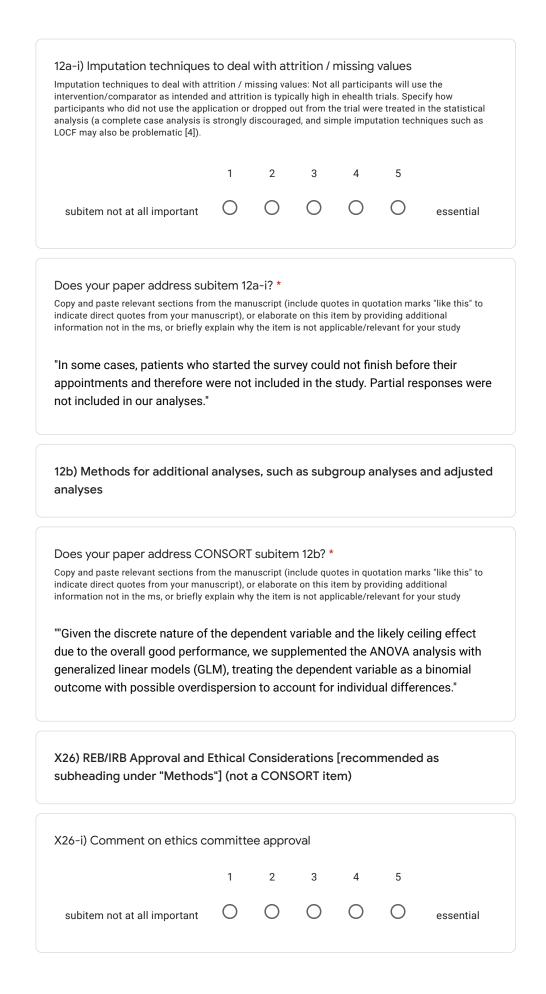
11a-i) Specify who was blinded, and who wasn't Specify who was blinded, and who wasn't. Usually, in web-based trials it is not possible to blind the participants [1, 3] (this should be clearly acknowledged), but it may be possible to blind outcome assessors, those doing data analysis or those administering co-interventions (if any).
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subitem not at all important
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Does your paper address subitem 11a-i? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"Responses were directly collected and stored on our server. Other than selfreported demographic information, no user information is stored on the server."





Does your paper address subitem X26-i?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"The work in this study was approved by the Institutional Review Boards (IRB) at the University of Massachusetts Medical School and LGH. All participants from LGH were shown an information sheet describing the study, had the ability to ask questions before participating, and provided verbal informed consent before participating. AMT participants provided electronic informed consent before participating."

x26-ii) Outline informed consent procedures Outline informed consent procedures e.g., if consent was obtained offline or online (how? Checkbox, etc.?), and what information was provided (see 4a-ii). See [6] for some items to be included in informed consent documents. 1 2 3 4 5 0 0 0 0 \bigcirc subitem not at all important essential

Does your paper address subitem X26-ii?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"All participants from LGH were shown an information sheet describing the study, had the ability to ask questions before participating, and provided verbal informed consent before participating. AMT participants provided electronic informed consent before participating."

X26-iii) Safety and security procedures

Safety and security procedures, incl. privacy considerations, and any steps taken to reduce the likelihood or detection of harm (e.g., education and training, availability of a hotline)

	1	2	3	4	5	
subitem not at all important	0	0	0	0	0	essential

Does your paper address subitem X26-iii?

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

13a) For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analysed for the primary outcome

NPT: The number of care providers or centers performing the intervention in each group and the number of patients treated by each care provider in each center

Does your paper address CONSORT subitem 13a? *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

"We recruited a total of 188 participants at the community hospital location from the end of December 2019 to the beginning of March 2020. Results from 174 participants were included in the final analysis. 141 recruited subjects were patients, 33 recruited subjects were persons accompanying patients."

13b) For each group, losses and exclusions after randomisation, together with reasons

Does your paper address CONSORT subitem 13b? (NOTE: Preferably, this is shown in a CONSORT flow diagram) *

Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study

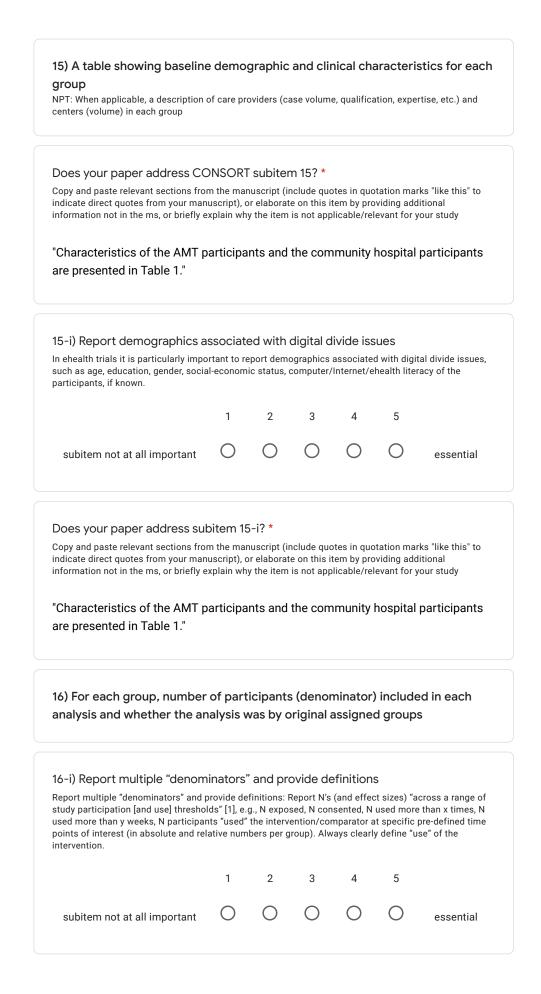
"14 participants were recruited and began the task, but did not complete it as they were called to their appointments and therefore were not included in our final analyses."

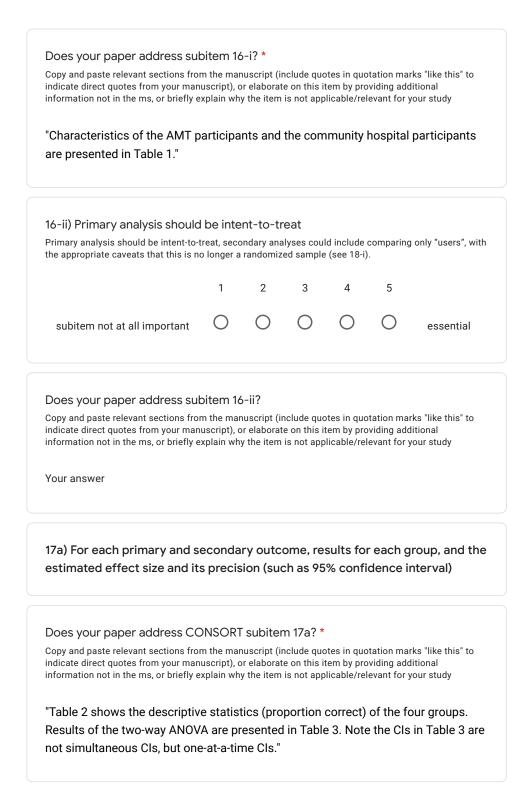
13b-i) Attrition diagram

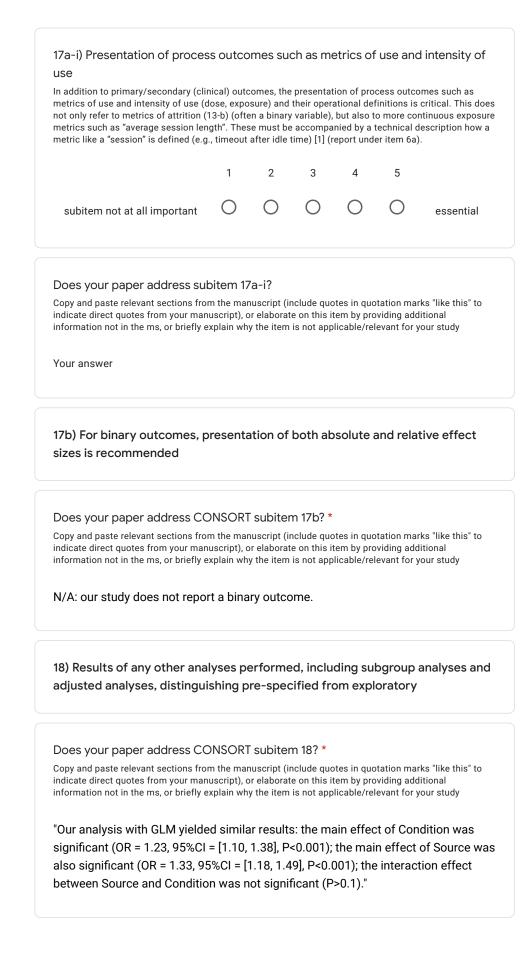
Strongly recommended: An attrition diagram (e.g., proportion of participants still logging in or using the intervention/comparator in each group plotted over time, similar to a survival curve) or other figures or tables demonstrating usage/dose/engagement.

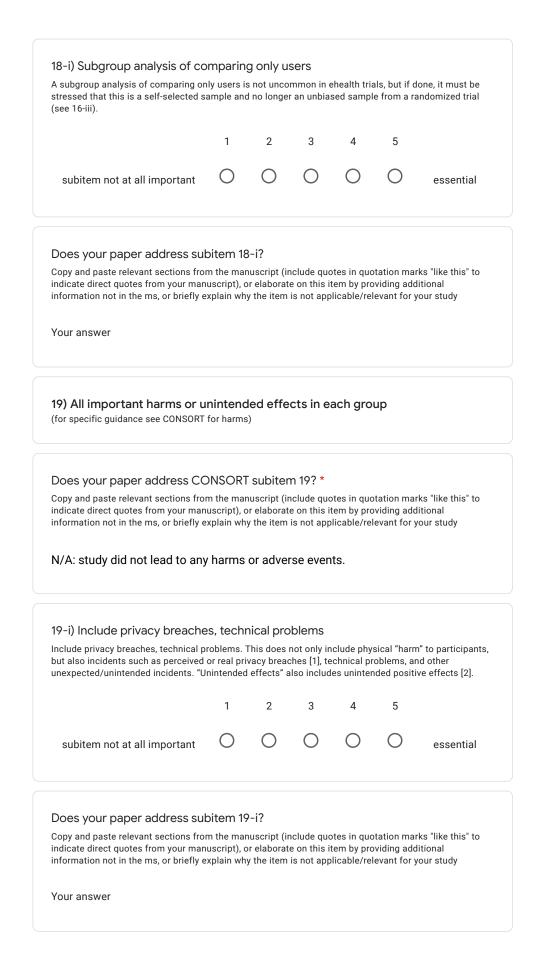
	1	2	3	4	5	
subitem not at all important	0	0	0	0	0	essential

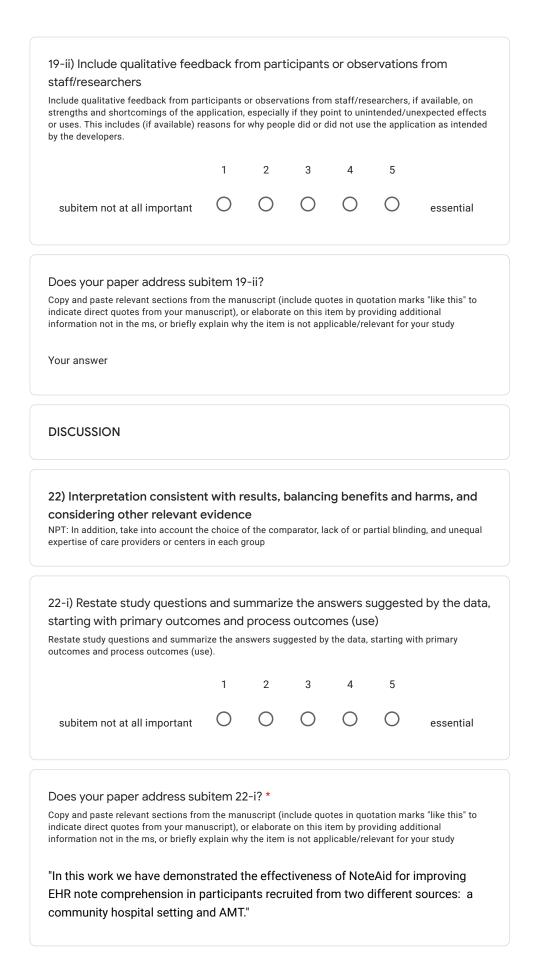
Does your paper address subitem 13b-i? Copy and paste relevant sections from the manuscript or cite the figure number if applicable (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study								
Your answer								
14a) Dates defining the periods of recruitment and follow-up								
Does your paper address CONSORT subitem 14a? * Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study "We recruited a total of 188 participants at the community hospital location from the end of December 2019 to the beginning of March 2020."								
14a-i) Indicate if critical "secular events" fell into the study period Indicate if critical "secular events" fell into the study period, e.g., significant changes in Internet resources available or "changes in computer hardware or Internet delivery resources"								
1 2 3 4 5 subitem not at all important O O O essential								
Does your paper address subitem 14a-i? Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study Your answer								
14b) Why the trial ended or was stopped (early)								
Does your paper address CONSORT subitem 14b? * Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study "local recruitment of individuals for our trial had to be halted due to COVID-19."								



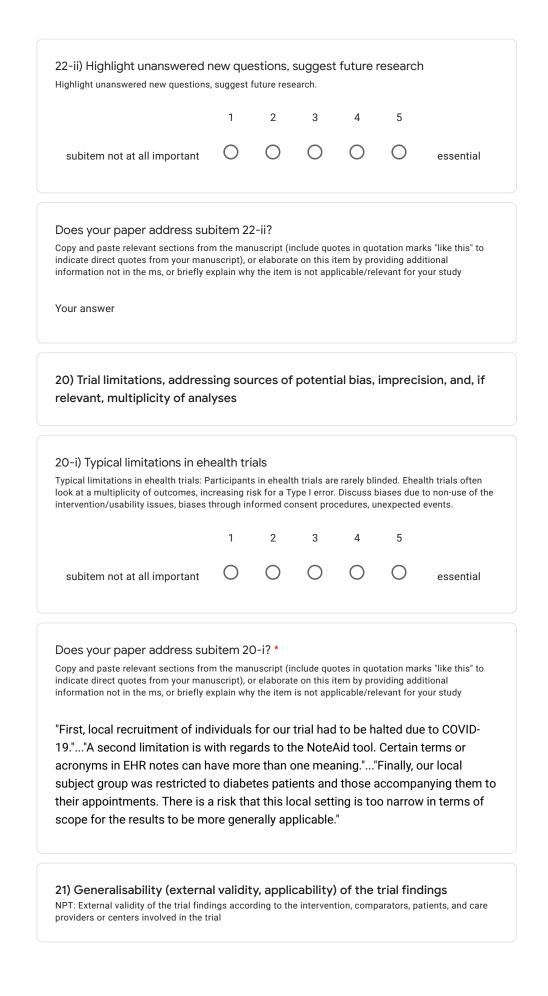




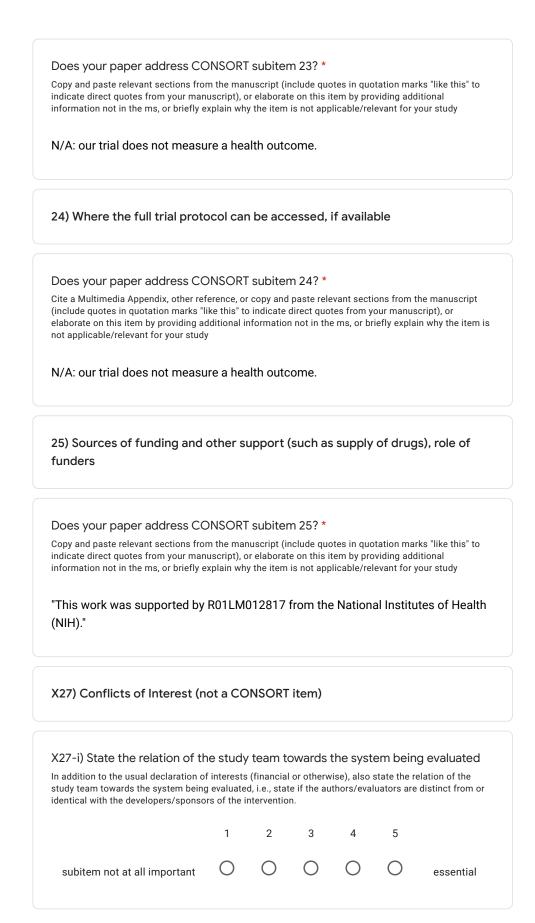




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21-ii) Discuss if there were e	lements	in the F	CT that	would l	oe differ	ent in a
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Does your paper address subitem X27-i? Copy and paste relevant sections from the manuscript (include quotes in quotation marks "like this" to indicate direct quotes from your manuscript), or elaborate on this item by providing additional information not in the ms, or briefly explain why the item is not applicable/relevant for your study "CONFLICTS OF INTEREST: None"
About the CONSORT EHEALTH checklist
As a result of using this checklist, did you make changes in your manuscript? *
yes, minor changes
What were the most important changes you made as a result of using this checklist? Your answer
How much time did you spend on going through the checklist INCLUDING making changes in your manuscript *
1-2 hours
As a result of using this checklist, do you think your manuscript has improved? *
() yes
○ no
O Other:

Would you like to become involved in the CONSORT EHEALTH group? This would involve for example becoming involved in participating in a workshop and writing an "Explanation and Elaboration" document	
() yes	
o no	
O Other:	
Clear select	ion
Any other comments or questions on CONSORT EHEALTH	
Your answer	
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