



5.1.2e

**Center for Infectious Disease Control | Epidemiology and surveillance unit | Department Enteric, Vectorborne and Zoonotic infections (EVZ)**

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**From:** 5.1.2e <5.1.2e@coneno.com>

**Sent:** donderdag 29 februari 2024 10:36

**To:** 5.1.2e <5.1.2e@rivm.nl>

**Cc:** 5.1.2e <5.1.2e@rivm.nl>; 5.1.2e <5.1.2e@rivm.nl>; 5.1.2e

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**Subject:** Re: CASE management with reworked auth - progress update

I think that would make sense to aim for. I will soon send technical details on how to start test deployment for this, so you can start collecting experience with the new tools and data flows in the next weeks.

If deployment goes well, this will simplify functional management of the system for sure,

It is also a step towards simplifying the backend architecture and includes better logging. As I understood, patientportal will need a pentest as well, so this might be included in the scope of that.

On 29. Feb 2024, at 10:09, 5.1.2e <5.1.2e@rivm.nl> wrote:

Looks and sounds great 5.1.2e

What do you think: can the implementation of LP-PLUS and the postcovidportal be a "test new-CASE" here, or better stay away from that?

Vriendelijke groeten,

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**From:** 5.1.2e <5.1.2e@coneno.com>  
**Sent:** donderdag 29 februari 2024 09:07  
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<5.1.2e@rivm.nl>; 5.1.2e <5.1.2e@rivm.nl>  
**Subject:** CASE management with reworked auth - progress update

Hi 5.1.2e

I hope you had a nice holiday.

In the last two weeks, we have been making excellent progress on the authorisation rework of the functional management. I attached my current task overview for this particular activity. (See attached PDF. For the colors: green is considered to be already implemented, yellow is a todo, red is a todo which likely will expand/broken down into further subtasks)

As you can see, a big part is already covered in green, and I would say these are also the parts that were core for the rework. For now, we work towards covering all the needed functionality of the system, so you can deploy and use it for all the tasks.

The core of the changes seems to work in my development version quite well:

- I can use my organisation's Azure AD (MS Entra ID) account to login if I am authorised by the organisations admin. By default, this would be a simple management-user account in CASE.
- In Azure AD, admins can assign the application specific role "ADMIN" to a user. Once this is done, after the next login, the CASE backend accepts this admin role, and considers the management user as an admin with access to all configs and user-management.

- management-users, who are not admins, need granted permissions for to access features of the management-functionality

- to authorise management-users, admins can create permissions:

- a permission contains: subjectID, subjectType, resourceType, resourceKey, action and optional limiters

- for example: a management user might have the permission: study - infectieradar - upload\_survey - limited to "weekly" → user can upload new versions of the weekly, but cannot do anything else not granted.

- admins can grant you the permission to perform all actions in all studies (study, \*, \*), or that you are also allowed to manage email templates, or other resources

- later we will share some documentation about it

All together this is a very big leap forward for the study software. The management UI will get improvements and features of the next weeks / months (in VERDI one goal is to have the improved survey editor, and study rule editor developed). But when the here mentioned changes will be in-place, this already makes the management of the system much more robust.

5.1.2e I will send infos later about steps on how to deploy the current versions of these.

Let me know, if you already have questions, we can schedule a call :-)

Best,

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<image001.png>

<image002.png>

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5.1.2e

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