

## SafeBoosC III Data completion report

25-May 2021

The data shown below is based on all registered data entries in OpenClinica on SafeBoosC III trial participants, up until 25th of May 2021. The data has been analysed and reported according to the SafeBoosC III Central monitoring plan (see [www.safeboosc.eu](http://www.safeboosc.eu) under “Good Clinical Practice”).

Direct link to the Central monitoring plan: <https://www.rigshospitalet.dk/english/departments/juliane-marie-centre/department-of-neonatology/research/SafeboosC-III/Documents/central-monitoring-plan.pdf>

### Completion of data entries

Below you will find an overview of data entries across all centres.

Note that completion calculations are based on babies that have reached at least 10 days of life for end-of-monitoring completion, and 40 weeks of postmenstrual age (PMA) for Follow-up and Blinded follow-up completion. n/a = not applicable, meaning that no babies at the specific site has reached 10 days of life or 40 weeks of PMA when applicable.

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
AT01 Univ. Hospital Graz	14	100%	100%	100%	63%
BE01 Univ. Hospital Leuven	25	100%	100%	90%	86%
BE02 AZ St. Jain Univ. Hosp. Bruges	10	100%	50%	100%	57%

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
BE03 Charleroi Univ. Hospital	6	100%	100%	80%	80%
BE04 CHU Tivoli Hospital	13	100%	100%	100%	67%
BE06 Liege Rocourt Hospital	16	56%	57%	100%	25%
CH01 University Hospital Zürich	37	100%	100%	100%	80%
CH03 University Hospital Lucern	31	100%	87%	100%	79%
CH04 University Hospital Geneva	8	100%	67%	100%	0%
CH05 Lausanne University Hospital	16	100%	86%	100%	0%
CN01 Children's Hospital, Zheijang Univ.	3	0%	0%	100%	0%
CN02 Children's Hospital, Fudan	35	100%	100%	94%	89%
CN04 Guangzhou Women and Children's Hospital	2	0%	0%	n/a	n/a
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
CN05 Longgang Distr. Centr. Hosp. Shenzhen	8	88%	100%	100%	100%
CN06 Xiamen Children's Hospital	4	100%	100%	100%	100%
CN07 The People's Hospital of Dehong	3	100%	100%	100%	33%
CN08 Maternal and Child Health Hosp. Quangxi	7	100%	100%	80%	80%
CZ01 The Institute for the Care of Mother and Child	49	100%	100%	100%	90%
CZ02 Motol Univ. Hospital	3	100%	n/a	100%	0%
DE01 University Hospital Freiburg	9	100%	100%	100%	100%
DK01 Rigshospitalet	82	99%	100%	100%	99%
DK04 Aalborg University Hospital	13	73%	57%	83%	92%
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
DK18 Aarhus University Hospital	13	100%	100%	100%%	56%
DK30 Odense University Hospital	9	44%	0%	33%	0%
ES01 La Paz University Hospital	65	100%	100%	100%	100%
ES02 Hospital Clinic de Barcelona (Maternitat)	45	100%	100%	100%	100%
ES03 University Hospital 12 de Octubre	50	100%	100%	100%	100%
ES05 Hospital de Sant Joan de Deu	25	100%	100%	100%	100%
ES06 H. U. Puerta del Mar	18	94%	88%	100%	100%
ES08 Hospital Clinico San Carlos	27	100%	100%	100%	100%
ES09 H. U. Marques de Valdecilla	16	94%	100%	79%	100%
ES10 Virgen de las Nieves	11	100%	100%	100%	100%
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
ES11 H. Univ. Juan XXIII Tarragona	14	79%	100%	100%	100%
ES13 Hospital de Cruces	2	100%	0%	n/a	n/a
GR01 Alexandra Hospital, Athens	13	100%	100%	100%	100%
GR02 Ippokrateion Hospital of Thessalonikki	28	100%	100%	100%	100%
GR03 Univ of Patras, General Hospital	9	100%	100%	100%	100%
GR04 Univ Hospital of Heraklion	8	100%	100%	100%	88%
IE01 Univ. College Cork	11	100%	100%	100%	0%
IE02 Rotunda Hospital Dublin	1	100%	n/a	n/a	n/a
IE04 NMH Holles St	1	n/a	n/a	n/a	n/a
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
IN01 St Johns Medical College Hospital, Bangalore	3	100%	100%	100%	100%
IT01 Presidio Ospedale Sant'Anna, Turin	5	100%	100%	100%	100%
IT07 Fondazione IRCCS Milano	33	97%	100%	100%	100%
IT08 Ospedale del Ponte, Varese	4	100%	100%	100%	100%
IT09 Fondaz. Policlinico Univ. A Gemelli	7	100%	100%	100%	100%
NY10 Oslo University Hospital	6	67%	0%	n/a	n/a
PL01 Medical Center UJASTEK Krakow	29	100%	100%	100%	91%
PL03 Specialist Hospital No. 2 Bytow	5	100%	0%	100%	100%
PL04 Poznan Univ. of Medical Sciences	27	89%	85%	100%	96%
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
PL07 Warsaw Univ. Medical Sciences	8	100%	100%	100%	75%
PL08 Szpital Uniwersytecki, Krakow	9	100%	100%	100%	100%
PL12 Centre of Medical Postgraduate Education, Warsaw	1	100%	0%	n/a	n/a
TR01 Gazi University Hospital	12	100%	100%	100%	88%
TR02 Marmara University Hospital	27	100%	100%	100%	100%
TR03 Uludag University Hospital	27	94%	100%	94%	94%
TR04 Kanuni Sultan Hospital	11	100%	100%	100%	100%
TR05 Bilkent, Ankara City Hospital	34	88%	92%	100%	64%
TR06 Basaksehir City Hospital	6	83%	0%	100%	100%
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Site	Randomisations up until 25-May 2021	End-of-monitoring (72 hrs) % completed	Serious adverse reactions (72 hrs) % completed	Follow-up (36 weeks) % completed	Blinded follow-up (brain ultrasound) % completed
UK08 Royal Hospital for Children, Glasgow	6	67%	100%	n/a	n/a
UK09 NHS Lanarkshire Hospital	4	100%	0%	n/a	n/a
US02 Loma Linda University Hospital	25	100%	100%	100%	57%
US03 University of Utah, Div. Neonatology	15	100%	100%	100%	100%
US04 UT Southwestern Medical Center	5	100%	100%	n/a	0%
<b>Total</b>	<b>1070</b>	<b>96%</b>	<b>93%</b>	<b>98%</b>	<b>86%</b>

Note that completion calculations are based on babies that have reached at least 10 days of life for end-of-monitoring completion, and 40 weeks of postmenstrual age (PMA) for Follow-up and Blinded follow-up completion. n/a = not applicable, meaning that no babies at the specific site has reached 10 days of life or 40 weeks of PMA when applicable.

Investigators who do not have 100% completion of data entries across all three modules will be contacted by e-mail and asked to enter the missing data into OpenClinica.

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