NGO Report to the 7th Periodic Report of France on the Convention against Torture (CAT)

+ Supplement “IGM Practices and the CAT”
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This NGO Report online:  
Executive Summary

France is in breach of its obligations under the Convention against Torture to (a) take effective legislative, administrative, judicial or other measures to prevent non-urgent, unnecessary surgery and other medical treatment carried out on intersex persons without the effective, informed consent of those concerned, causing severe mental and physical pain and suffering, and (b) to ensure impartial investigation, access to redress, and the right to fair and adequate compensation and rehabilitation for victims. (Arts. 2, 12, 14 and 16, General Comments 2 and 3). (A, B, C)

This Committee has already recognised such non-consensual, unnecessary treatments as a breach of the Convention in previous Concluding Observations for Germany, Switzerland, Austria, Denmark and Hong Kong, and called for legislation to (a) end the practice and (b) ensure redress and compensation, plus (c) access to free counselling. In addition, also CRC, CRPD, CEDAW, the UN Special Rapporteur on Torture (SRT), the UN High Commissioner for Human Rights (UNHCHR), the World Health Organisation (WHO) and the Council of Europe (COE) have all repeatedly recognised IGM Practices as a serious human rights violation, and have called for legislative remedy (SRT, CRC, COE) and for access to redress and justice for victims (CRC, CRPD, WHO) (F, Annex 2).

Intersex people are born with Variations of Sex Anatomy, including atypical genitals, atypical sex hormone producing organs, atypical response to sex hormones, atypical genetic make-up, atypical secondary sex markers. While intersex people may face several problems, in the “developed world” the most pressing are the ongoing Intersex Genital Mutilations, which present a distinct and unique issue constituting significant human rights violations (D).

IGM Practices include non-consensual, medically unnecessary, irreversible, cosmetic genital surgeries, and/or other harmful medical treatments that would not be considered for “normal” children, without evidence of benefit for the children concerned, but justified by societal and cultural norms and beliefs. (E.1.) Typical forms of IGM include “masculinising” and “feminising”, “corrective” genital surgery, sterilising procedures, imposition of hormones, forced genital exams, vaginal dilations, medical display, human experimentation and denial of needed health care (E.2., “IGM in Medical Textbooks”).

IGM Practices cause known lifelong severe physical and mental pain and suffering, including loss or impairment of sexual sensation, painful scarring, painful intercourse, incontinence, urethral strictures, impairment or loss of reproductive capabilities, lifelong dependency of artificial hormones, significantly elevated rates of self-harming behaviour and suicidal tendencies, lifelong mental suffering and trauma, increased sexual anxieties, less sexual activity, dissatisfaction with functional and aesthetic results. (E, Cases No. 1–2)

Since 1950, IGM has been practised systematically and on an industrial scale all over the “developed world”, and all typical IGM forms are still practised in France today. Parents and children are misinformed, kept in the dark, sworn to secrecy, kept isolated and denied appropriate support. (A, E, Cases No. 1–2, “IGM in Medical Textbooks”).

For more than 20 years, intersex people, NGOs, human rights and bioethics bodies have criticised IGM as harmful and traumatizing, as a fundamental human rights violation, as a form of genital mutilation and child sexual abuse, as torture or ill-treatment, and called for legislation to end it and to ensure remedies (F).

This Thematic NGO Report has been compiled by French intersex person and advocate Vincent Guillot and the international intersex NGO Zwischengeschlecht.org / StopIGM.org. It contains Concluding Recommendations (C).
NGO Report
to the 7th Periodic Report of France
on the Convention against Torture (CAT)

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Introduction

Background and State Report

France will be considered for its seventh periodic review by the Committee against Torture in its 57th Session in 2016. In France, doctors in public, university and private clinics are regularly performing IGM practices, i.e. non-consensual, medically unnecessary, irreversible cosmetic genital surgeries, sterilising procedures, and other harmful treatments on intersex children, which have been described by survivors as genital mutilation and torture. IGM practices are known to cause severe, lifelong physical and psychological pain and suffering, and have been repeatedly recognised by this Committee and other UN bodies as constituting torture or ill-treatment, violence and a harmful practice.

Unfortunately, human rights violations of intersex children and adults weren’t mentioned in the State Report. However, this NGO Report demonstrates that the current medical treatment of intersex infants and children in France constitutes a serious breach of France’s obligations under the Convention on the Rights of the Child.

The French State not only does nothing to prevent this abuse, but in fact directly finances it via the public health assurances and via funding the public university clinics and paediatric hospitals, thus violating its duty to prevent harmful practices. To this day the French Government refuses to take appropriate legislative, administrative and other measures to protect intersex children, and refuses survivors the right to justice, redress and compensation.

About the Rapporteurs

This NGO report has been prepared by the French intersex person and advocate Vincent Guillot in collaboration with the international intersex NGO Zwischengeschlecht.org / StopIGM.org.

- **Vincent Guillot** is a French intersex person, survivor of IGM practices and intersex human rights defender who has been working to improve the well-being and human rights of intersex persons in France and Europe, and to raise awareness on intersex issues for more than a decade. Vincent is a co-founder of Organisation Intersex International (OII).

- **Zwischengeschlecht.org / StopIGM.org**, founded in 2007, is an international Human Rights NGO based in Switzerland. It is led by intersex persons, their partners, families and friends, and works to represent the interests of intersex people and their relatives, raise awareness, and fight IGM Practices and other human rights violations perpetrated on intersex people, according to its motto, “Human Rights for Hermaphrodites, too!” According to its charter, Zwischengeschlecht.org works to support persons concerned seeking redress and justice, and has continuously collaborated with members of parliament and other bodies in order to call on Governments and Clinics to collect and disclose statistics of intersex births and IGM practices, and to prevent them.

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1. [http://www.histoiresordinaires.fr/Intersexe-Vincent-Guillot-sort-de-la-nuit_a1330.html](http://www.histoiresordinaires.fr/Intersexe-Vincent-Guillot-sort-de-la-nuit_a1330.html)
4. [https://www.woz.ch/-2044](https://www.woz.ch/-2044)
Methodology

This thematic NGO report is a localised and updated addition to the 2015 thematic CAT NGO Reports for Switzerland, Austria and Denmark by partly the same rapporteurs.4

This Report includes two anonymised case studies of French survivors of IGM practices. The stories were originally obtained for the 2015 CRC NGO Report by the same Rapporteurs,5 the identity of the witnesses being known to the Rapporteurs. Each first-person narrative is preceded by a standardised abstract composed by the Rapporteurs. The small number of case studies is due to the fact that many patients, their families, and parents find it hard to speak about what happened to them, and do not wish their story to become public, even anonymously. These cases, however, show in an exemplary manner how IGM are practiced without informed consent by the persons concerned and/or their parents, and cause severe pain and suffering, both physical and psychological.

Intersex Genital Mutilations are still an “emerging human rights issue,” unfortunately often neglected due to lack of access to comprehensive information. To assess the current practice at national level, some general knowledge on the matter is crucial. For further reference, and to facilitate access to more comprehensive information for the Committee, the rapporteurs attached abbreviated thematic supplements.6

The rapporteurs are aware that IGM practices are a global issue, which can’t be solved on a national level alone. However, this report illustrates why France is a State Party to which it would be timely and most appropriate to issue strong recommendations.

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A. IGM Practices in France

1. Lack of Protection for Intersex Persons, IGM Practices Remain Pervasive

a) Overview

In France (see CRC/C/FRA/CO/5, paras 47-48), same as in the neighbouring states of Switzerland (see CAT/C/CHE/CO/7, para 20; CRC/C/CHE/CO/2-4, paras 42-43) and Germany (see CAT/C/DEU/CO/5; para 20; CRPD/C/DEU/CO/1, paras 37-38), there are no legal or other protections in place to ensure the rights of intersex children to physical and mental integrity, autonomy and self-determination, and to prevent non-consensual, medically unnecessary, irreversible surgery and other harmful treatments a.k.a. IGM practices.

To this day, the French government, despite now acknowledging the “extremely recent issue”, undeviatingly refuses to “take effective legislative, administrative, judicial or other measures” to protect intersex children, but instead claims the “expertise” of the perpetrators themselves would be best suited to “prevent” practices “that later may be experienced as intolerable mutilations” (see below A.2.4. “Lack of legislative provisions”).

At the same time, all forms of IGM practices remain widespread and ongoing, advocated by the official public medical body “Haute Autorité de Santé (HAS)”, including in “National Guidelines”, and paid for by the public health insurance.

Currently, all major French public University or Regional Children’s Clinics (including, but not limited to the government-appointed “Reference centres for rare diseases of sex development”) employ doctors advocating, prescribing and perpetrating IGM practices.

b) Most Common IGM Forms advocated by Officially Appointed Bodies and Doctors

• IGM 3: Sterilising Procedures plus arbitrary imposition of hormones (see also Cases No. 1–2), as advocated by Prof. Pierre Mouriquand, chief surgeon of the government-appointed “Reference centres for rare diseases of sex development”:

  “Surgery on the gonads
  • Removal of the gonads in CAIS and PAIS because of risk of tumours in adults.
  • Early surgery or surgery during puberty, according to the choice of the parents:
    • Associated inguinal hernia
    • Psychological problems according to age”

See also photos of gonadectomies by Pierre Mouriquand on p. 61

• IGM 2: Feminising Genital Surgeries, as advocated by the official public medical body “Haute Autorité de Santé (HAS)” in their “National CAH Guidelines”:

  “4.4 Surgical Therapy
  4.4.1 Environment
  “[...] The surgical treatment is prescribed by the paediatric surgeon according to surgery for anomalies of sex development.

  4.4.2 Surgical Schedule
  “French surgeons operate on the little girls when the metabolic and endocrine situation is stable, earliest in the first months of life. The essential reasons for choosing this age is the responsiveness of genital

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tissues when the repair is done early, and the minimisation of psychological consequences for the child and the parents."

**4.4.5 Surgical Procedure**

"The surgical procedure during the first months of life includes three principal stages:

- opening of the vaginal cavity at the pelvic floor (vaginoplasty), which represents the most difficult part, in particular in cases of high confluence
- if necessary, the reduction in size of the clitoris while preserving the vascularisation and the nerves
- the perineoplasty which, if possible, consists of the reconstruction of the small labia, the margins of the vaginal introitus, and the reduction of the labia majora which are often enlarged."

See also photo of “feminising labioplasty” by Pierre Mouriquand on p. 60

- **IGM 1: Masculinising Genital Surgeries** (see also Case No. 1), as advocated by the official public medical body “Haute Autorité de Santé (HAS)”:

  "1.2.3. Position of the working group
  
  […] According to the working group, [penile] curvatures associated with a hypospadias […] are ideally operated in early infancy […]."  

According to a publication by a public medical body under the auspices of the French Ministry of Health and based on statistics obtained from the official hospital activity and expenditure data ("Programme de médicalisation des systèmes d’information (PMSI)"), in France over 3000 children under 7 years are submitted to “hypospadias repair surgery” every year, with the majority of children under 2 years at the time of surgery, and with total numbers increasing yearly, and the age at surgery becoming lower and lower.  

See also photos of “masculinising hypospadias repair” by Pierre Mouriquand on p. 59

- **Repeated Forced Genital Exams and Photography** (see also Case No. 1) are also common place in France, see e.g. the pictorial examples published by the “French Urology Association”.  

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c) Examples of French University Children’s Clinics advocating & perpetrating IGM

- **Centre Hospitalier Universitaire de Lyon**, government-appointed “Reference centre for rare diseases of sex development”, see various pictorial and text examples in publications and presentations. Prof. Pierre Mouriquand and fellow surgeon Dr. Daniela Gorduza are also part of the surgeon’s working group responsible for the “National CAH Guidelines” advocating “feminising” IGM practices (see above). CHU de Lyon is also part of the international perpetrator’s study “DSDnet”, where surgeon Pierre Mouriquand is listed as participant, as well as part of the international perpetrator’s study “DSD-Life”, where surgeon Daniela Gorduza is listed in the project staff (for background information on both “DSDnet” and “DSD-Life” see also below p. 46–47).

Prof. Mouriquand is also known as a particularly adamant advocate of IGM practices consciously dismissing human rights concerns (see below p. A.1.d).

See also pictorial examples of IGM practices by Pierre Mouriquand on p. 59–61

- **Hôpitaux universitaires Paris-Sud (Hôpital Kremlin-Bicêtre)**, government-appointed “Reference centre for rare diseases of sex development”, is also part of the international perpetrator’s study “DSD-Life” (see also below p. 46), where endocrinologist Dr. Claire Bouvattier is listed as the scientific lead, and the centre (self-)described as follows:

> “The Department of Pediatric Endocrinology, Hôpital Kremlin Bicêtre, has a longstanding experience in clinical care of patients with DSD. Dr. Claire Bouvattier sees 250-300 patients/year with DSD in their clinics and have access to all DSD patients of associated hospitals through the French National DSD Reference centre. The French National DSD Reference Paris centre comprises a multidisciplinary specific DSD team including: Pediatric Endocrinology, Pediatric Radiology, Hormonal and Molecular Biology [sic], Pediatric Surgery, Gynecology and Psychology [sic]. [...] As our DSD center is a partner for all the French pediatric endocrinologists, we will coordinate the study DSD-Life in France.”

A 2014 “DSD-Life”-presentation co-authored by Dr. Claire Bouvattier lists for “Individuals with DSD” the following common surgical procedures:

- See e.g. Pierre Mouriquand: “Prise en charge chirurgicale des Anomalies Congénitales du Développement Génito-Sexuel (DSD): La féminisation” (2015), [online]
- Pierre Mouriquand: “Anomalies congénitales du développement génito-sexuel (DSD).” (2012), [online]
And an online fact sheet “Doctor, is it a boy or a girl?” co-authored by Dr. Claire Bouvattier and paediatric surgeon Dr. Daniela Gordoza (CHU Lyon, see above) claims:

“(12) The goal of the surgical intervention is to give back [sic] an anatomy and functionality of the genital apparatus by carrying out feminising or masculinising surgery.” 18

• Hôpital Universitaire mère-enfant Robert-Debré, Paris lists as “Specialty: surgery on anomalies of the genitourinary system” 19 Prof. Alaa El-Ghoneimi, consulting surgeon for paediatric urology, is also part of the surgeon’s working group of the French National CAH guidelines (see above A.1.b “feminising surgeries”), and a particularly adamant advocate of IGM practices consciously dismissing human rights concerns (see below A.1.d).

• Lille University Clinic (Hôpital Jeanne de Flandre), in a 2015 presentation by paediatric surgeons of the clinic, counts “1003 cases of hypospadias were identified in our reference center” from 1992–2012. 20

• Amiens University Clinic, in a 2015 presentation by paediatric surgeons of the clinic, counts 57 cases of hypospadias including 10.5% cases of undescended testes enrolled in a 2011–2014 study. 21

d) French IGM Doctors consciously dismissing Human Rights Concerns

It must be duly noted that French paediatric surgeons are particularly adamant advocates of IGM practices, consciously dismissing to consider any human rights concerns, despite openly acknowledging knowledge of relevant criticisms by human rights and ethics bodies.

For example Prof. Alaa El-Ghoneimi (Hôpital Universitaire mère-enfant Robert-Debré, Paris, see above) simply dismissed the 2013 Report by the Special Rapporteur on Torture as “unjust”. 22
In the same vein, Prof. Pierre Mouriquand (Centre Hospitalier Universitaire de Lyon, see above) dismissed both the 2013 Report by the Special Rapporteur on Torture and the 2012 Recommendations by the Swiss National Advisory Commission on Biomedical Ethics as “inappropriate and biased statements” and “biased and counterproductive reports”, while insisting on continuing with IGM practices.23

2. The Treatment of Intersex Persons in France as Torture

a) Infliction of Severe Pain or Suffering

It is well established that IGM Practices generally inflict lifelong, severe pain and suffering (see p. 50–53). Cases No. 1–2 prove in an exemplary manner that this is also true in France.

b) Intention

It is generally established that surgery on intersex persons is always intentionally performed and not merely the result of negligence, and that it does not detract from the intention if doctors perform surgery for well-meant purposes, see p. 53–54. Cases No. 1–2 prove that this is also true in France.

c) Purpose of Discrimination

It is generally established that on the basis of their “indeterminate sex,” intersex children are singled out for experimental harmful treatments that would be “considered inhumane” on “normal” children. Thus intersex children are penalised compared to “normal” infants, even where the perpetrator has benign intentions, see p. 54. The evidence from French clinics, and medical publications and guidelines prove this also to be true in France, as do Cases No. 1–2.

d) Involvement of a State Official

In France with its public and mandatory health assurances paying for the medical ill-treatment of intersex persons and its government-appointed “Reference centres for rare diseases of sex development” and “National Guidelines” advocating and perpetrating IGM, it is self-evident that, even if IGM practices would take place in a Private Clinic, it is directly attributable to the state, and was committed at the very least with the acquiescence of a person acting in an official capacity; and even more so in the case of public University Clinics, Federal State Clinics and government-appointed institutions. As is the failure of the State to exercise due diligence to protect this group of citizens from torture.

e) Lawful Sanction

Non-consensual unnecessary surgery performed on an intersex child or adult does not constitute a sanction in France. It is therefore not covered by the exception clause.

3. The Treatment of Intersex Persons in France as Ill-Treatment

Even if it would be considered that the treatment of intersex people in France does not constitute torture, it certainly constitutes cruel, inhuman and degrading treatment (Art. 16, see p. 55). Ill-treatment is equally prohibited by the Convention in absolute and non-derogable terms. According to the Committee’s General Comment 3, for CIDT also Article 14 applies.

4. Lack of Legislative Provisions, Impunity of the Perpetrators

Art. 2 of the Convention obliges State parties to “take effective legislative, administrative, judicial or other measures to prevent acts of torture in any territory under its jurisdiction.” General Comment 2 states, “The obligation to prevent ill-treatment in practice overlaps with and is largely congruent with the obligation to prevent torture,” and similarly obliges State parties to “to eliminate any legal or other obstacles that impede the eradication of torture and ill-treatment; and to take positive effective measures to ensure that such conduct and any recurrences thereof are effectively prevented.”

Accordingly, with regards to IGM practices, the Committee already explicitly recognised the obligation for State parties to “Take the necessary legislative, administrative and other measures to guarantee respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected during infancy or childhood to non-urgent medical or surgical procedures.”

However, the French State party, despite now admitting to being made aware of the “extremely recent issue,” and having tasked the National Consultative Ethics Committee for health and life sciences (Comité consultatif national d’éthique pour les sciences de la vie et de la santé) to report on the matter, so far still refuses to “take effective legislative, administrative, judicial or other measures” to protect intersex children, but instead claims the “expertise” of the perpetrators themselves would be best suited to ensure prevention of further serious human rights violations of intersex children:

“With such complex medical questions, which entail multiple lifelong consequences, the quality of the medical expertise and care, and of a continuous dialogue is best suited to guarantee compliance with the rights of the child. It must ensure to prevent any premature decisions and any action, which later may be experienced as intolerable mutilation.”

5. Obstacles to Redress, Fair and Adequate Compensation

The statutes of limitation prevent survivors of early childhood IGM Practices to call a court because persons concerned often do not find out about their medical history until

24 Committee against Torture (2008), General comment No. 2, CAT/C/GC/2, para. 3-4.
28 Current Minister of Family, Children and Women’s Rights, Mrs Laurence Rossignol in the French Senate on the topic of prevention of IGM practices (10.02.2016) [own translation], http://stop.genitalmutilation.org/post/France-Gov-claims-perpetrators-best-suited-to-prevent-IGM.
29 Ibid.
much later in life, which in combination with severe trauma caused by IGM Practices often proves to amount to a severe obstacle. Even though in France the statutes of limitations compare favourably to neighbouring countries, they still effectively prohibit survivors of early childhood IGM practices to call a court, as persons concerned often do not find out about their medical history until much later in life, and severe trauma caused by IGM Practices often prohibits them to act in time once they do. So far there was no case of a victim of IGM practices succeeding in going to court (see also Case No. 2).

The French government so far refuses to ensure that non-consensual unnecessary IGM surgeries on minors are recognised as a form of torture or ill-treatment (see above), or as a form of genital mutilation or harmful practice respectively, which would case formally prohibit parents from giving “consent”. In addition, the State party refuses to initiate impartial investigations, as well as data collection, monitoring, and disinterested research (see below p. 46). In addition, despite legal obligations to do so, also French hospitals are often unwilling to provide full access to patient’s files (see also Cases No. 1–2).

This situation is not in line with France’s obligations under the Convention.

B. Conclusion: France is Failing its Obligations towards Intersex People under the Convention against Torture

The surgeries and other harmful treatments intersex people endure cause severe physical and mental pain and suffering. Doctors perform the surgery for the discriminatory purpose of making a child fit into societal and cultural norms and beliefs, although there is plenty of evidence on the suffering this causes. The State party is responsible for these violations amounting to torture or at least ill-treatment, committed by publicly funded doctors, clinics, and universities, as well as in private clinics, all relying on money from the mandatory health insurance, and public grants. Although meanwhile the pervasiveness IGM practices is common knowledge, and the State party even admits to being made aware of the “extremely recent issue”, France nonetheless fails to prevent these grave violations both in public and in private settings, but allows the human rights violations of intersex children, adolescents and adults to continue unhindered.

Thus France is in breach of its obligation to take effective legislative, administrative, judicial or other measures to prevent acts of torture (Art. 2 CAT) or other forms of cruel, inhuman or degrading treatment (Art. 16 CAT, General Comment 2).

Also in France, victims of IGM practices encounter severe obstacles in the pursuit of their right to an impartial investigation (Arts. 12, 13 CAT), and to redress and fair and adequate compensation, including the means for as full rehabilitation as possible (Art. 14 CAT, General Comment 3).

Also the State party’s efforts on education and information regarding the prohibition against torture in the training of medical personnel are grossly insufficient with respect to the treatment of intersex people (Art. 10 CAT).

30 Globally, no survivor of early surgeries ever managed to have their case heard in court. All relevant court cases (3 in Germany, 1 in the USA) were either about surgery of adults, or initiated by foster parents.

C. Recommendations

The Rapporteurs respectfully suggest that the Committee recommends the following measures to the French Government with respect to the treatment of intersex Persons (based on the Committee’s previous recommendations to Switzerland, Austria and Denmark):

Intersex Persons

The Committee welcomes the Government’s decision to task the National Consultative Ethics Committee for health and life sciences (Comité consultatif national d’éthique pour les sciences de la vie et de la santé) to report on unnecessary surgery and other medical treatment carried out on intersex persons (i.e. persons with variations of sex anatomy) without the effective, informed consent of those concerned.

The Committee remains concerned, however, that these procedures, which reportedly caused physical and psychological suffering, have not as yet given rise to any inquiry, sanction or reparation (arts. 2, 12, 14 and 16).

The Committee recommends that the State party:

(a) Take the necessary legislative, administrative and other measures to guarantee the respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected to unnecessary medical or surgical procedures during infancy or childhood, but that all non-urgent medical interventions are postponed until a child is sufficiently mature to participate in decision-making and give full, free and informed consent;

(b) Guarantee impartial counselling services and psychosocial support for all intersex children and their parents, so as to inform them of the consequences of non-urgent, unnecessary surgery and other medical treatment and the possibility of postponing any decision on such treatment or surgery until the persons concerned are able to decide for themselves;

(c) Undertake investigation of reports of surgical and other medical treatment of intersex people without effective consent and adopt legal provisions in order to provide redress to the victims of such treatment, including adequate compensation.
Annexe 1 “Case Studies”

The first-person narratives have been collected for this thematic NGO report. The abstracts were composed by the Rapporteurs. The identity of all persons concerned is known to the Rapporteurs.

Case Study No. 1

The person concerned was born 1965 in the Clinique des Sablons in Neuilly sur Seine, with a micro penis, without scrotum and testes, and a vaginal cavity. The diagnosis was “déficit gonadique primitif” / “anorchia” (absence of testes, gonadal dysgenesis, streak gonads). The child was raised as a boy. The gonads were removed, and a scar on the penis suggests a hypospadias surgery. A test at age 10 showed male chromosomes XY. Later a gynaecologist confirmed the probable removal of an uterus. The person suffers to this day from the treatment and lies, and is unable to work.

The person concerned tells their story:

I was born 1965 in the Clinique des Sablons in Neuilly sur Seine, with a micropenis, without scrotum and testes, and with a hidden vaginal cavity, as the third of six children, to a professor of history and a nurse. I was hospitalised immediately at the Hôpital Saint-Vincent-de-Paul in Paris for fifteen days. Neither me nor my parents know what happened during these days. When I was returned to my parents, they were only informed that I had suffered an allergic reaction to antibiotics. I was eventually diagnosed with „déficit gonadique primitif“ (gonadal dysgenesis) and „anorchia“ (absence of testes). When my mother picked me up, the doctors told her that I was a monster and that I would die soon.

Until the age of six I lived a normal life with my parents and siblings, and went to school regularly. Nobody told me anything and I never asked questions.

At six years, I underwent a first exploratory surgery without results, performed by doctor Ficheux at the Clinique Villa Medicis in Courbevoie, which was declared as appendectomy. I have no idea if they already removed my gonads or if they did it later.

After this surgery my parents bought a house in the Bretagne, twelve hours from our home, where I would live most of the time during the year, together with my grandmother and a schizophrenic uncle, because my parents were ashamed of me, especially after the surgeries.

After having raised thirteen children herself, my grandmother didn’t want me. She was always very severe and ordered me around. Even when I was week after surgery, I had to make my bed alone. I had to help a lot on my grandmother’s farm, went to school rarely and didn’t have much contact with peers. When my grandmother had to attend a family celebration, my uncle was put in a hospital and I was sent to stay with another family. At the funeral of my mother a year ago, a lot of my relatives were surprised to see me, they thought I had died as a child.

Although at the age of seven my life was normal, and I didn’t have any pain or difficulties urinating, until the age of majority I was treated quarterly at the Hôpital Trousseau in Paris by urologist Jacques Brueziere, head of department Béatrice Mougenot, and endocrinologist M.C. Raux-Eurin.

I only stayed with my parents when I had to go to the hospital in Paris for my quarterly examinations or for surgeries. During that time I also attended school near Paris. My siblings and classmates never understood why I was absent so often. When I was well I would stay at
my parent’s house for a month or so. When I had surgery they would send me back to my grandmother in the Bretagne.

During the quarterly appointments at the hospital, where I had to stay for some days or even a week, the doctors performed X-rays to determine the bone maturity, they took blood samples, urinary probes, they examined my naked body, looked at my genitals, checked the growth of my armpit hair, and took photographs. Afterwards, they first talked with my parents alone, then with me and my parents together. I also had to see a doctor who performed psychological tests, but didn’t even know he was a psychologist.

I was intimidated and afraid. The doctors never told me or my parents anything but, “Everything is OK”, or “don’t cry, you are big”. They told me that there is nobody else like me, that I am the only one on this planet.

I suffered very early of anxiety, and our family doctor prescribed me tranquillisers.

The doctors told my parents that I was a failed boy, but they would fix me, and that everything was alright. They regularly gave me testosterone injections.

Since my mother couldn’t get any answers out of the doctors, she asked a personal friend who happened to be a clinician herself. I know from correspondence in my medical records that both, my mother’s doctor friend and our family doctor, were instructed to tell us exactly the same: that everything is OK, I just had to take the testosterone and attend the quarterly examinations. My mother was very disappointed when she found out.

In the following years I underwent another three large surgeries at the Hôpital Trousseau in Paris, which also are referred to as functional explorations in my medical records without further specifications. My parents weren’t informed about the results, or what was done exactly.

A total of ten scars on my abdomen suggest even more surgeries which aren’t mentioned in the medical records I was eventually able to obtain.

After surgery the doctors would always apply an urethral catheter, although I was able to go to the toilet myself – unlike other children in the same room, which for example had a broken leg, but used a urine bottle.

My urethral catheter was often removed roughly, although I told the nurse I felt it was sticking, which was very painful. I regularly ended up with having infections, blood and pus in my urine, which required a treatment with antibiotics.

During the morning visits, often 10–15 medical personnel and students came into the room, the Professor lifted my blanket, and everybody stared at my genitals.

My father used to bring me to hospital, and my mother came to take me home afterwards. But they both never visited me in the hospital. My father said he was too busy, and my mother couldn’t bear it. Once she had encountered me lying in bed, hands and feet bound, and was shocked. Only two young aunts came to see me once a week. Sometimes it was a friend of my parents, who brought me to the hospital. When my mother came to the hospital to take me home, we didn’t take a taxi, but went with the bus to the train station, although I couldn’t barely stand straight because I was weak and in pain. My mother was ashamed, because people looked at me, and told me to pull myself together.

My mother told me more than once it would have been better, if I was born dead. „If I had known, I would have aborted you.”

At thirteen, the doctors prescribed me regular shots of testosterone. Nonetheless, I developed
breasts during puberty which I hid under baggy clothes. I hated the testosterone shots, they
gave me headaches and tremendous pain in my legs, I was aggressive and couldn’t sleep, nor
concentrate at school. However, the doctors always blamed something else, preferably grow-
ing pains or my attitude, and insisted on continuing with the shots. Eventually, at sixteen I
simply refused to see them anymore, and until the age of 22 I had no hormones at all.

I found out myself that I had a vaginal cavity which was closed by a membrane. When I was
22 a gynaecologist confirmed that I have a small vaginal cavity. Some years ago another gyn-
aecologist told me that based on the informations in the medical records, I must have had at
least the rudiment of an uterus, which obviously had been removed.

I was able to obtain some of my medical records in 2003 with the help of my former family
doctor, who told me that, because of a new law in France from 2002, every patient would
have the right to have his medical records. So he ordered my medical records and we studied
them together. They weren’t complete, the years until I turned six were missing, and there
were some sheets which obviously belonged to another boy’s records.

This was the first time I learned that I am intersex. I was happy, because it proved I wasn’t
crazy after all.

Nonetheless, in my adult years my bad experiences with doctors and hospitals continued.
Several times I was forced to or blackmailed into “consenting“ to hormonal treatments or
surgeries, for example when my parents admitted me to a clinic after a suicide attempt.

I have been taking anxiolytics, antidepressants and mood stabilisers from the age of eighteen.
When I stopped with the anxiolytics for a few months, my genital and my legs started to hurt.
My doctor told me that the anxiolytics were also painkillers and had suppressed the pain.

To this day I can’t sleep except when I am dead tired or with the aid of barbiturates, because
I have nightmares either of surgeries or medical exams, or training courses for assistant doc-
tors, where they displayed me naked.

I still suffer from an intense pain when I urinate, recurring urinary tract infections, and in-
continence.

Today I refuse any follow-up treatments, and any hormone replacement therapy with testos-
sterone, but I take estrogen without prescription instead. My body didn’t change. However, I
no longer suffer from hot flashes, and I could quit the mood-stabilisers which I was prescribed
for many years.

Since 2006, I am unable to work due to anxiety and incontinence. At 50, I never made any
contribution to a retirement pension, I am without home, and on welfare.
Case Study No. 2

The person concerned was born 1971 in Nantes and raised as a girl. When turning 17, she consulted different doctors due to absent puberty, without letting her parents know. It was found that she had XY chromosomes and mixed gonads, but the doctors withheld the truth from her. She submitted to gonadectomy without informed consent, but was given conflicting justifications for the surgery by different doctors. 17 years later, after years of denial and breakdown, the person concerned returned to her former surgeon and finally get’s to know the truth.

The person concerned tells their story:

In 1988, at the age of 17, after waiting to hit puberty in vain for years, I decided to consult a gynaecologist. I chose a specialist in Nantes, about 50 kilometres far from where I lived, instead of going to the family doctor, the subject being tabu at home. I left home hitchhiking, pretending to go to an appointment with the nephrologist who for years has been seeing me regularly for a renal insufficiency. I was welcomed by the gynaecologist Dr Françoise Dano who assured me that everything was normal. She prescribed me a blood test. When I met her again, her tone had changed. She was less confident and sent me to an endocrinologist. Therefore, I found myself as a sick person for something that I didn’t link with health.

The endocrinologist, Dr Hubert du Rostu, prescribed me a karyotype test. This examination reminded me of biology course at school, where we studied the human karyotype, showing but one anomaly, trisomy 21. I therefore asked myself what had brought them to suspect me in such a way. I went to see Dr Dano, the gynaecologist, to know about the result of this examination. She put down the document in front of me and explained that there was no anomaly. I couldn’t stop looking at the paper where I read that my sex chromosomes were XY, which seemed to contradict my biology course. My knowledge vanished in front of the confidence with which Dr Dano assured me that everything was OK.

I returned to high school the next day and couldn’t help myself to brag about having had a chromosomal analysis which turned out to be perfect, thus XY. I was so happy that they hadn’t found trisomy. The entire class burst out laughing, and I was told that this was impossible, since XY are male sex chromosomes. I was confused and told myself that I might have read it wrong. I still was waiting for an injection or a treatment which would induce my puberty, the only reason why I had turned to a doctor. Instead of that, the endocrinologist and the gynaecologist decided to send me to the Brétésché clinic in Nantes to see the urologist and surgeon Dr Éric Hermouët to have my ovaries examined with a camera and under general anaesthetic. I left the clinic without knowing more about it.

I saw Dr Dano again with the idea that everything was finally settled. She told me again that they had to observe the ovaries with the naked eye via exploratory surgery before the treatment could start. I was hospitalised for two and a half days for this intervention. I was asked to enter the clinic on a Sunday evening, Dr Hermouët received me in the deserted outpatients department, and made me sign a consent form allowing him to remove everything which had to be. I was underage then. I didn’t meet Dr Hermouët after the surgery and couldn’t get any information before leaving the clinic. I went to see the family doctor a week later to remove the stitches like the nurse at the clinic had asked me.

I encountered his substitute who took note of the letter from the surgeon in front of me. He seemed stunned. When he realised I hadn’t been informed at all, his jaw dropped, and he eventually told me that they had removed my ovaries. This was the first blow. I realized that I would never have children. The substitute recommended to return to see the family doctor after he’d be back from his holiday. I went to see Dr André Le Cardiec the next week. However,
I didn’t encounter the warm person who had cared for all my family, but an outraged man who was shouting at me. As I showed no reaction to his words which I couldn’t comprehend, he completely lost his temper and told me that my kidney failure was something rare, but what they had just discovered inside of me was extremely rare and monstrous. I was shocked by his agitation and his offending words.

They had discovered that I was a monster, my life left me and endless emptiness took its place. I walked out of the doctor’s office watching the sky hoping that someone over there come and tell me that I had landed on the wrong planet. I didn’t know where to go. Then I decided to go back to my mother’s place and I entered into a period of self-destruction which lasted two years. Two weeks after this horrifying news, I saw Dr Dano who softened the subject talking about a removal of ovaries that had been necessary to avoid cancer. She assured me that I was a complete woman and that it would be possible for me to bear children with hormonal therapy and an egg cell donation. I clung to this hope and entered into adulthood with the hormonal treatment. Then I cut off with all these doctors except Dr Dano, the gynaecologist. Two years later, in 1990, I couldn’t remember what had happened.

17 years later, after years of denial and breakdowns, I decided to meet Dr Hermouët again to understand the nature of the surgery I was submitted to in 1988. First he denied and explained that it was his father, which had retired, who had operated me. Then he excused himself and asked me to leave his office because he had to take care of a child birth. I realized that he tried to hide something and I waited for him to return in front of his door. He returned half an hour later, holding my medical records in his hands. He then invited me to enter his study and warned me that what was inside of this file was unspeakable. I insisted to know and he explained that my karyotype turned out to be male, and that they had removed one gonad on the occasion of the first surgical examination, which turned out to be a testicle. The second had been removed during the second surgery.

Anyway, that was different from what I was told earlier. I was both proud and shaky when I visited Dr Dano, the gynaecologist who meanwhile had refused to let me try assisted reproduction with egg cell donation. I told her what I had learned. She categorically denied everything, called me delirious and suggested to call a psychiatrist for an appointment. Also when I tried to talk to her about this subject later, she always insisted she wasn’t aware of this and that she wasn’t my doctor at that time. When I asked her why she prescribed me hormones, she answered that I suffered from early menopause or from ovarian cancer, insisting that this was the only information given to her. Then I found some support in a community center, before meeting other intersex persons via OII Francophonie.

In 2015, I tried to get my medical records back from Dr du Rostu’s office. As soon as he received my letter he called me back claiming that he had no file with my name in his office. I then threatened to come with a lawyer which gave him a scare. He promised to look for it. A few hours later he called me back to say he couldn’t find anything with my name in his records. A week later, I received a one page document that was difficult to read. It was the result of the analysis of the first testicle removed, describing ovarian tissue without egg cells and showing no risk of malignancy. A week later I received three more pages of my records among which there was the analysis of the second gonad describing a mixture of ovarian and testicular tissues, but still with no risk of malignancy. Therefore, Dr Hermouët had lied to me when he claimed that he had found testicles instead of ovaries, to avoid me to contest these operations. Between 2007 and 2015, the statutes of limitations which are twenty years in France had expired and it was too late to register a complaint.
Annexe 2 “Bibliography: IGM in Human Rights Mechanisms”

1. International Bodies Recognising Human Rights Violations of Intersex Persons

2006: UN WHO, Genomic resource centre, Gender and Genetics: Genetic Components of Sex and Gender (online)

Gender Assignment of Intersex Infants and Children

Intersex is defined as a congenital anomaly of the reproductive and sexual system. An estimate about the birth prevalence of intersex is difficult to make because there are no concrete parameters to the definition of intersex. The Intersex Initiative, a North-American based organization, estimates that one in 2,000 children, or five children per day in the United States, are born visibly intersex. (36) This estimate sits within range from genital anomalies, such as hypospadias, with a birth prevalence of around 1:300 to complex genital anomalies in which sex assignment is difficult, with a birth prevalence of about 1:4500. (37) Many intersex children have undergone medical intervention for health reasons as well as for sociological and ideological reasons. An important consideration with respect to sex assignment is the ethics of surgically altering the genitalia of intersex children to “normalize” them.

Clitoral surgery for intersex conditions was promoted by Hugh Hampton Young in the United States in the late 1930s. Subsequently, a standardized intersex management strategy was developed by psychologists at Johns Hopkins University (USA) based on the idea that infants are gender neutral at birth. (38) Minto et al. note that “the theory of psychosexual neutrality at birth has now been replaced by a model of complex interaction between prenatal and postnatal factors that lead to the development of gender and, later, sexual identity.” (39) However, currently in the United States and many Western European countries, the most likely clinical recommendation to the parents of intersex infants is to raise them as females, often involving surgery to feminize the appearance of the genitalia. (40)

Minto et al. conducted a study aiming to assess the effects of feminizing intersex surgery on adult sexual function in individuals with ambiguous genitalia. As part of this study, they noted a number of ethical issues in relation to this surgery, including that:

- there is no evidence that feminizing genital surgery leads to improved psychosocial outcomes;
- feminizing genital surgery cannot guarantee that adult gender identity will develop as female; and that
- adult sexual function might be altered by removal of clitoral or phallic tissue. (41)

2009: UN CEDAW, CEDAW/C/DEU/CO/6, 10 February 2009, para 61–62:
http://www2.ohchr.org/english/bodies/cedaw/docs/co/CEDAW-C-DEU-CO6.pdf

Cooperation with non-governmental organizations

61. [...] The Committee regrets, however, that the call for dialogue by non-governmental organizations of intersexual [...] people has not been favourably entertained by the State party.

62. The Committee request the State party to enter into dialogue with non-governmental organizations of intersexual [...] people in order to better understand their claims and to take effective action to protect their human rights.

Follow-up to concluding observations

67. The Committee requests the State party to provide, within two years, written information on the steps undertaken to implement the recommendations contained in paragraphs 40 and 62.

2009: UN SR Health, A/64/472, 10 August 2009, para 49:
http://www.refworld.org/pdfid/4aa762e30.pdf

IV. Vulnerable groups and informed consent

A. Children

49. Health-care providers should strive to postpone non-emergency invasive and irreversible interventions until the child is sufficiently mature to provide informed consent. [67] [Fn. 67: This is particularly problematic in the case of intersex genital surgery, which is a painful and high-risk procedure with no proven medical benefits; see, e.g., Colombian Constitutional Court, Sentencia SU-337/99 and Sentencia T-551/99.] Safeguards should be in place to protect children from parents withholding consent for a necessary emergency procedure.
2011: UNHCHR, A/HRC/19/41, 17 November 2011, para 57:

“In addition, intersex children, who are born with atypical sex characteristics, are often subjected to discrimination and medically unnecessary surgery, performed without their informed consent, or that of their parents, in an attempt to fix their sex.”

2011: UN CAT, CAT/C/DEU/CO/5, 12 December 2011, para 20:
http://www2.ohchr.org/english/bodies/cat/docs/co/CAT.C.DEU.CO.5_en.pdf

**Intersex people**

20. The Committee takes note of the information received during the dialogue that the Ethical Council has undertaken to review the reported practices of routine surgical alterations in children born with sexual organs that are not readily categorized as male or female, also called intersex persons, with a view to evaluating and possibly changing current practice. However, the Committee remains concerned at cases where gonads have been removed and cosmetic surgeries on reproductive organs have been performed that entail lifelong hormonal medication, without effective, informed consent of the concerned individuals or their legal guardians, where neither investigation, nor measures of redress have been introduced. The Committee remains further concerned at the lack of legal provisions providing redress and compensation in such cases (arts. 2, 10, 12, 14 and 16).

The Committee recommends that the State party:

(a) Ensure the effective application of legal and medical standards following the best practices of granting informed consent to medical and surgical treatment of intersex people, including full information, orally and in writing, on the suggested treatment, its justification and alternatives;

(b) Undertake investigation of incidents of surgical and other medical treatment of intersex people without effective consent and adopt legal provisions in order to provide redress to the victims of such treatment, including adequate compensation;

(c) Educate and train medical and psychological professionals on the range of sexual, and related biological and physical, diversity; and

(d) Properly inform patients and their parents of the consequences of unnecessary surgical and other medical interventions for intersex people.

2013: UN SR Torture, A/HRC/22/53, 1 February 2013, paras 77, 76, 88:

77. Children who are born with atypical sex characteristics are often subject to irreversible sex assignment, involuntary sterilization, involuntary genital normalizing surgery, performed without their informed consent, or that of their parents, “in an attempt to fix their sex”, [107] leaving them with permanent, irreversible infertility and causing severe mental suffering.

76. [...] These procedures [genital-normalizing surgeries] are rarely medically necessary,[106] can cause scarring, loss of sexual sensation, pain, incontinence and lifelong depression and have also been criticized as being unscientific, potentially harmful and contributing to stigma (A/HRC/14/20, para. 23). [...]  

88. The Special Rapporteur calls upon all States to repeal any law allowing intrusive and irreversible treatments, including forced genital-normalizing surgery, involuntary sterilization, unethical experimentation, medical display, “reparative therapies” or “conversion therapies”, when enforced or administered without the free and informed consent of the person concerned. He also calls upon them to outlaw forced or coerced sterilization in all circumstances and provide special protection to individuals belonging to marginalized groups.

2013: Council of Europe (COE), Resolution 1952 (2013) “Children’s right to physical integrity”, 1 October 2013, paras 2, 6, 7:

2. The Parliamentary Assembly is particularly worried about a category of violation of the physical integrity of children, which supporters of the procedures tend to present as beneficial to the children themselves despite clear evidence to the
contrary. This includes, amongst others, female genital mutilation, the circumcision of young boys for religious reasons, early childhood medical interventions in the case of intersex children and the submission to or coercion of children into piercings, tattoos or plastic surgery.

6. The Assembly strongly recommends that member States promote further awareness in their societies of the potential risks that some of the above mentioned procedures may have on children’s physical and mental health, and take legislative and policy measures that help reinforce child protection in this context.

7. The Assembly therefore calls on member States to:

7.1. examine the prevalence of different categories of non-medically justified operations and interventions impacting on the physical integrity of children in their respective countries, as well as the specific practices related to them, and to carefully consider them in light of the best interests of the child in order to define specific lines of action for each of them;

7.2. initiate focused awareness-raising measures for each of these categories of violation of the physical integrity of children, to be carried out in the specific contexts where information may best be conveyed to families, such as the medical sector (hospitals and individual practitioners), schools, religious communities or service providers; [...]  

7.4. initiate a public debate, including intercultural and interreligious dialogue, aimed at reaching a large consensus on the rights of children to protection against violations of their physical integrity according to human rights standards;

7.5. take the following measures with regard to specific categories of violation of children’s physical integrity: [...]  

7.5.3. undertake further research to increase knowledge about the specific situation of intersex people, ensure that no-one is subjected to unnecessary medical or surgical treatment that is cosmetic rather than vital for health during infancy or childhood, guarantee bodily integrity, autonomy and self-determination to persons concerned, and provide families with intersex children with adequate counselling and support; [...]  

7.7. raise awareness about the need to ensure the participation of children in decisions concerning their physical integrity wherever appropriate and possible, and to adopt specific legal provisions to ensure that certain operations and practices will not be carried out before a child is old enough to be consulted.

2014: UN CRPD, CRPD/C/DEU/Q/1, 17 April 2014, paras 12–13:  
http://tbinternet.ohchr.org/Treaties/CRPD/Shared%20Documents/DEU/CRPD_C-DEU_Q_1_17084_E.doc

Freedom from exploitation, violence and abuse (art. 16)

12. How many irreversible surgical procedures have been undertaken on intersexual children before an age at which they are able to provide informed consent? Does the State party plan to stop this practice?

13. Please provide up to date statistics on forced sterilizations of persons, i.e. without their free and informed consent.

2014: WHO, OHCHR, UN Women, UNAIDS, UNDP, UNFPA, and UNICEF, Eliminating forced, coercive and otherwise involuntary sterilization. An interagency statement, May 2014, p 2, 6, 7:  
http://www.who.int/iris/bitstream/10665/112848/1/9789241507325_eng.pdf?ua=1

Background

Some groups, such as […] intersex persons, also have a long history of discrimination and abuse related to sterilization, which continues to this day. […] Intersex persons, in particular, have been subjected to cosmetic and other non-medically necessary surgery in infancy, leading to sterility, without informed consent of either the person in question or their parents or guardians. Such practices have also been recognized as human rights violations by international human rights bodies and national courts (15, 64).

[...] [I]ntersex persons

Intersex persons may be involuntarily subjected to so-called sex-normalizing or other procedures as infants or during childhood, which, in some cases, may result in the termination of all or some of their reproductive capacity. Children who are born with atypical sex characteristics are often subjected to cosmetic and other non-medically indicated surgeries performed on their reproductive organs, without their informed consent or that of their parents, and without taking into consideration the views of the children involved (64; 147, para 57; 148; 149). As a result, such children are being subjected to irreversible interventions that have lifelong consequence for their physical and mental health (64; 150, para
Medical procedures that might result in sterility may sometimes be justified because of benefits to health, including the reduction of cancer risk (152). Such treatments may be recommended for [...] intersex persons; however, they may be proposed on the basis of weak evidence, without discussing alternative solutions that would retain the ability to procreate (151, 153–157). Parents often consent to surgery on behalf of their intersex children, including in circumstances where full information is lacking (151, 158, 159).

It has been recommended by human rights bodies, professional organizations and ethical bodies that full, free and informed consent should be ensured in connection with medical and surgical treatments for intersex persons (64, 150) and, if possible, irreversible invasive medical interventions should be postponed until a child is sufficiently mature to make an informed decision, so that they can participate in decision-making and give full, free and informed consent (15, 149). It has also been recommended that health-care professionals should be educated and trained about bodily diversity as well as sexual and related biological and physical diversity, and that professionals should properly inform patients and their parents of the consequences of surgical and other medical interventions (149; 150, para 20; 160–162).

Remedies and redress

- Recognize past or present policies, patterns or practices of coercive sterilization, and issue statements of regret or apology to victims, as components of the right to remedy for these practices.
- Provide notification, through appropriate and humane means, to people who have been subjected to coercive sterilization, and who may be unaware of their situation, and provide information on the possibility of seeking administrative and judicial redress.
- Promptly, independently and impartially investigate all incidents of forced sterilization with due process guarantees for the alleged suspect, and ensure appropriate sanctions where responsibility has been established.
- Provide access, including through legal aid, to administrative and judicial redress mechanisms, remedies and reparations for all people who were subjected to forced, coercive or involuntary sterilization procedures, including compensation for the consequences and acknowledgement by governments and other responsible authorities of wrongs committed. Enable adults to seek redress for interventions to which they were subjected as children or infants.
- Guarantee access to reversal procedures, where possible, or assisted reproductive technologies for individuals who were subjected to forced, coercive or otherwise involuntary sterilization.

Monitoring and compliance

- Establish monitoring mechanisms for the prevention and documentation of forced, coercive and otherwise involuntary sterilization, and for the adoption of corrective policy and practice measures.
- Collect data regarding forced, coercive and otherwise involuntary sterilization, in order to assess the magnitude of the problem, identify which groups of people may be affected, and conduct a comprehensive situation and legal analysis.
- Providers of sterilization services should implement quality improvement programmes to ensure that recommendations aimed at preventing forced, coercive and otherwise involuntary sterilization are followed and procedures are properly documented.
- Establish mechanisms for obtaining patient feedback on the quality of services received, including from marginalized populations.


D. Violence against children (arts. 19, 24, para. 3, 28, para. 2, 34, 37 (a) and 39) [...] Harmful practices

42. While welcoming the adoption of a new provision of criminal law prohibiting genital mutilation, the Committee is deeply concerned at: [...] (b) Cases of medically unnecessary surgical and other procedures on intersex children, which often entail irreversible consequences and can cause severe physical and psychological suffering, without their informed consent, and the lack of redress and compensation in such cases.

43. The Committee draws the attention of the State party to the Joint General Comment
No. 18 on harmful practices (2014), together with the Committee on the Elimination of Discrimination against Women, and urges the State party to: [...] 
(b) In line with the recommendations on ethical issues relating to intersexuality by the National Advisory Commission on Biomedical Ethics, ensure that no-one is subjected to unnecessary medical or surgical treatment during infancy or childhood, guarantee bodily integrity, autonomy and self-determination to children concerned, and provide families with intersex children with adequate counselling and support.

2015: UN CRPD, CRPD/C/DEU/CO/1, 13 May 2015, p. 6–7, paras 37-38: 

Protecting the integrity of the person (art. 17)

37. The Committee is concerned about: [...] c) the lack of implementation of the 2011 recommendations CAT/C/DEU/CO/5, para. 20, regarding upholding bodily integrity of intersex children.

38. The Committee recommends that the State party take the necessary measures, including of a legislative nature to:

[...]

(d) Implement all the recommendations of CAT/C/DEU/CO/5, para. 20 relevant to intersex children.

https://wcd.coe.int/ViewDoc.jsp?Ref=CommDH/IssuePaper%282015%291&Language=lanEnglish&Ver=original&BackColorInternet=C3C3C3&BackColorIntranet=EDB021&BackColorLogged=F5D383

http://apps.who.int/iris/bitstream/10665/175556/1/9789241564984_eng.pdf?ua=1

“A major concern for intersex people is that so-called sex normalizing procedures are often undertaken during their infancy and childhood, to alter their bodies, particularly the sexual organs, to make them conform to gendered physical norms, including through repeated surgeries, hormonal interventions and other measures. As a result, such children may be subjected to medically unnecessary, often irreversible, interventions that may have lifelong consequences for their physical and mental health, including irreversible termination of all or some of their reproductive and sexual capacity.”

“Increasingly, concerns are being raised by intersex people, their caregivers, medical professionals and human rights bodies that these interventions often take place without the informed consent of the children involved and/or without even seeking the informed consent of their parents (178, 262, 264, 270–273).”

“It has also been recommended [by human rights bodies and ethical and health professional organizations] that investigation should be undertaken into incidents of surgical and other medical treatment of intersex people without informed consent and that legal provisions should be adopted in order to provide remedies and redress to the victims of such treatment, including adequate compensation (91, 264).”

2015: UN CAT, CAT/C/CHE/CO/7, 14 August 2015, para 20: 

Intersex persons

20. The Committee welcomes the Federal Council decision to give an opinion by the end of 2015 on the recommendations of the National Advisory Commission on Biomedical Ethics with regard to the unnecessary and in some cases irreversible surgical procedures that have been carried out on intersex persons (i.e. persons with variations in sexual anatomy) without the effective, informed consent of those concerned. However, the Committee notes with concern that these procedures, which reportedly caused physical and psychological suffering, have not as yet given rise to any inquiry,
sanction or reparation (arts. 2, 12, 14 and 16).

The Committee recommends that, in light of the forthcoming decision by the Federal Council, the State party:

a) Take the necessary legislative, administrative and other measures to guarantee respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected during infancy or childhood to non-urgent medical or surgical procedures intended to decide the sex of the child, as recommended by the National Advisory Commission on Biomedical Ethics and the Committee on the Rights of the Child (see CRC/C/CHE/CO/2-4, para. 43 (b));

b) Guarantee counselling services and free psychosocial support for all persons concerned and their parents, and inform them that any decision on unnecessary treatment can be put off until the person concerned are able to decide for themselves;

c) Undertake investigation of reports of surgical and other medical treatment of intersex people without effective consent and adopt legal provisions in order to provide redress to the victims of such treatment, including adequate compensation.

2015: UN CRC, CRC/C/CHL/CO/4-5, 2 October 2015, paras 48–49:

Harmful practices

48. While noting the proposed development of a protocol for the health care of intersex babies and children, the Committee is seriously concerned about cases of medically unnecessary and irreversible surgery and other treatment on intersex children, without their informed consent, which can cause severe suffering, and the lack of redress and compensation in such cases.

49. In the light of its joint general comment No. 18 (2014) and No. 31 of the Committee on the Elimination of Discrimination against Women on harmful practices, the Committee recommends that the State party expedite the development and implementation of a rights-based health care protocol for intersex children, setting the procedures and steps to be followed by health teams, ensuring that no one is subjected to unnecessary surgery or treatment during infancy or childhood, protecting the rights of the children concerned to physical and mental integrity, autonomy and self-determination, providing intersex children and their families with adequate counselling and support, including from peers, and ensuring effective remedy for victims, including redress and compensation.

2015: UN CAT, CAT/C/AUT/CO/6, 9 December 2015, paras 44–45:

Intersex Persons

44. The Committee appreciates the assurances provided by the delegation that surgical interventions on intersex children are carried out only when necessary, following medical and psychological opinions. It remains concerned however about reports on cases of unnecessary surgery and other medical treatment with life-long consequences to which intersex children would have been subjected without their informed consent. The Committee is further concerned at the lack of legal provisions providing redress and rehabilitation in such cases (arts. 14 and 16).

45. The State party should:

(a) Take the necessary legislative, administrative and other measures to guarantee the respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected during infancy or childhood to non-urgent medical or surgical procedures intended to decide the sex of the child;

(b) Guarantee impartial counselling services for all intersex children and their parents, so as to inform them of the consequences of unnecessary and non-urgent surgery
and other medical treatment to decide on the sex of the child and the possibility of postponing any decision on such treatment or surgery until the persons concerned can decide by themselves;

(c) Guarantee that full, free and informed consent is ensured in connection with medical and surgical treatments for intersex persons and that non-urgent, irreversible medical interventions are postponed until a child is sufficiently mature to participate in decision-making and give effective consent;

(d) Undertake investigation of instances of surgical interventions or other medical procedures performed on intersex people without effective consent, and ensure that the persons concerned are adequately compensated.

2015: UN CAT, CAT/C/DNK/CO/6-7, 9 December, paras 42–43:

Intersex Persons

42. While taking note of the information provided by the delegation on the decision-making process related to treatment of intersex children, the Committee remains concerned at reports of unnecessary and irreversible surgery and other medical treatment with life-long consequences to which intersex children have been subjected before the age of 15 when their informed consent is required. The Committee is further concerned at hurdles faced by these persons when seeking redress and compensation in such cases (arts. 14 and 16).

43. The State party should:

(a) Take the necessary legislative, administrative and other measures to guarantee the respect for the physical integrity and autonomy of intersex persons and ensure that no one is subjected during infancy or childhood to unnecessary medical or surgical procedures;

(b) Guarantee counselling services for all intersex children and their parents, so as to inform them of the consequences of unnecessary surgery and other medical treatment;

(c) Ensure that full, free and informed consent is respected in connection with medical and surgical treatments for intersex persons and that non-urgent, irreversible medical interventions are postponed until a child is sufficiently mature to participate in decision-making and give full, free and informed consent;

(d) Provide adequate redress for the physical and psychological suffering caused by such practices to intersex persons.

2015: UN CAT, CAT/C/CHN-HKG/CO/4-5, 9 December 2015, paras 28–29:

[...][I]ntersex Persons

28. [...] The Committee is also concerned that intersex children are subjected to unnecessary and irreversible surgery to determine their sex at an early stage. Furthermore, the Committee is concerned at the long term physical and psychological suffering caused by such practices (arts. 10, 12, 14 and 16).

29. HKSAR should:

[...]

(b) Guarantee impartial counselling services for all intersex children and their parents, so as to inform them of the consequences of unnecessary and non-urgent surgery and other medical treatment to decide on the sex of the child and the possibility of postponing any decision on such treatment or surgery until the persons concerned can decide by themselves;

(c) Guarantee that full, free and informed consent is ensured in connection with medical and surgical treatments for intersex persons and that non-urgent, irreversible medical interventions are postponed until a child is sufficiently mature to participate
in decision-making and give full, free and informed consent;
    
    (d) Provide adequate redress for the physical and psychological suffering
cau sed by such practices to some intersex persons.

2016 UN CRC, CRC/C/IRL/CO/3-4, 4 February 2016, paras 39-40:
http://stop.genitalmutilation.org/public/Ireland_Concl-Obs_CRC_C_IRL_CO_3-4_22988_E.doc

E. Violence against children (arts. 19, 24, para.3, 28, para. 2, 34, 37 (a) and 39)
Harmful practices [CRC art. 24(3)]

39. The Committee notes as positive the adoption of the Gender Recognition Act of 2015 by the State party. How-
ever, the Committee remains concerned about cases of medically unnecessary surgeries and other procedures on intersex
children before they are able to provide their informed consent, which often entail irreversible consequences and can cause
severe physical and psychological suffering, and the lack of redress and compensation in such cases

40. The Committee recommends that the State party:

(a) Ensure that no-one is subjected to unnecessary medical or surgical treatment dur-
    ing infancy or childhood, guarantee bodily integrity, autonomy and self-determination
to children concerned, and provide families with intersex children with adequate
counselling and support;

(b) Undertake investigation of incidents of surgical and other medical treatment of
    intersex children without informed consent and adopt legal provisions in order to pro-
    vide redress to the victims of such treatment, including adequate compensation; and,

(c) Educate and train medical and psychological professionals on the range of sexual,
    and related biological and physical, diversity and on the consequences of unnecessary
    surgical and other medical interventions for intersex children.

2016 UN CRC, CRC/C/FRA/CO/5, 4 February 2016, paras 47-48:
http://stop.genitalmutilation.org/public/France_Concl-Obs_CRC_C_FRA_CO_5_22995_E.doc

D. Violence against children (arts. 19, 24, para.3, 28, para. 2, 34, 37 (a) and 39)
Harmful practices [CRC art. 24(3)]

47. While noting with appreciation the progress made by the State party to eradicate female genital mutilation the
Committee is, however, concerned by many young girls still at risk and the possible resurgence of the phenomenon.
The Committee is also concerned that medically unnecessary and irreversible surgery and other treatment is routinely
performed on intersex children.

48. Recalling the Committee’s joint general comment No. 18 (2014) and No. 31 of the Com-
mittee on the Elimination of Discrimination against Women on harmful practices, the
Committee recommends that the State party gather data with a view to understanding the
extent of these harmful practices with a view to better identify children at risk and pre-
vent them. It recommends that the State party:

(a) Increase awareness of female genital mutilation in the State party among girls
    at risk, medical professionals, social workers, the police and gendarme officers, and
    magistrates;

(b) Develop and implement a rights-based health care protocol for intersex children,
    ensuring that children and their parents are appropriately informed of all options,
    that children are involved, to the largest extent, in decision-making about their treat-
    ment and care, and no child is subjected to unnecessary surgery or treatment.
2. State Bodies Recognising Human Rights Violations of Intersex Persons


2013: Australian Senate, Community Affairs References Committee, Involuntary or coerced sterilisation of intersex people in Australia, October 2013

2014: German Conference of Women’s and Equality Ministers (GFMK), Resolution of the 24th GFMK Conference, 1–2 October 2013

2015: Maltese Parliament, Gender Identity Gender Expression and Sex Characteristics Act (GIGESC), 14 April 2015, Article 14(1–5) “Right to bodily integrity and physical autonomy”

2015: Austrian Children’s and Youth Attorneys (KiJAÖ) [NHRI], Position Paper on Intersex, [7 October 2015]
http://kija.at/images/KiJAOE-Positionspapier_zur_Intersexualitt_2015.pdf

3. National Ethics Bodies Recognising Human Rights Violations of Intersex Persons

2011: German Ethics Council, Opinion Intersexuality, 23 February 2012

2012: Swiss National Advisory Commission on Biomedical Ethics (NEK-CNE), On the management of differences of sex development. Ethical issues relating to “intersexuality”, Opinion No. 20/2012, 9 November 2012

4. NGO, NHRI Reports on Human Rights Violations of Intersex Persons

2004: CESCR Argentina, Mauro Cabral

2008: CEDAW Germany, Intersexuelle Menschen e.V./XY-Frauen

2010: CEDAW Germany, Intersexuelle Menschen e.V./XY-Frauen

2011: CEDAW Costa Rica, IGLHRC / MULABI, p. 8–11

2011: CAT Germany, Intersexuelle Menschen e.V./XY-Frauen, Humboldt Law Clinic
2012: UPR Switzerland, Swiss NGO Coalition for the UPR, para 18
http://lib.ohchr.org/HRBodies/UPR/Documents/Session14/CH/JS3_UPR_CHE_S14_2012_JointSubmission3_E.pdf

2012: UN SRT, Advocates for Informed Choice (AIC),

2012: CRC Luxemburg, Radelux


2013: CRPD Germany, BRK-Allianz, Germany, p. 36–37
http://www.brk-allianz.de/attachments/article/93/Alternative_Report_German_CRPD_Alliance_final.pdf

2013: UPR Germany, German Institute for Human Rights (GIHR), para 23
- German CRPD ALLIANCE, para 15
http://lib.ohchr.org/HRBodies/UPR/Documents/Session16/DE/js4_upr16_deu_s16_2013_jointsubmission4_e.pdf
- National Coalition for the Implementation of the UN Convention on the Rights of the Child in Germany (NC), para 4
http://lib.ohchr.org/HRBodies/UPR/Documents/Session16/DE/js5_upr_deu_s16_2013_jointsubmission5_e.pdf
- Forum Menschenrechte, paras 38, 39, 58
http://lib.ohchr.org/HRBodies/UPR/Documents/Session16/DE/js6_upr_deu_s16_2013_jointsubmission6_e.pdf

2013: CRC Germany, German Institute for Human Rights (GIHR), para 2.b.
- National Coalition for the Implementation of the UN Convention on the Rights of the Child in Germany (NC), lines 789–791, 826–828
http://www2.ohchr.org/english/bodies/crc/docs/ngos/Germany_National%20Coalition%20for%20the%20Implementation%20of%20the%20UNCRC%20in%20Germany_CRC%20Report-CRCWG65.pdf

2013: Inter-American Commission on Human Rights, Advocates for Informed Choice (AIC)
+ Hearing

2014: UNHRC, Canadian HIV/AIDS Legal Network, joined by ILGA

2014: CRC Switzerland, Child Rights Network Switzerland, p. 25–26
2014: CRC Switzerland, Zwischengeschlecht.org, Intersex.ch, SI Selbsthilfe Intersexualität

2014: CAT Australia, OII Australia, AISSGA, People with Disabilities, National LGBTI Health Alliance

2014: CAT USA, Advocates for informed Choice (AIC)

2015: CAT New Zealand, ITANZ

2015: CRPD Germany, Zwischengeschlecht.org

2015: CAT Switzerland, Zwischengeschlecht.org

2015: CRC Chile, Observatorio de Derechos Humanos – Chile (Andrés Rivera Duarte), International Gay and Lesbian Human Rights Commission (IGLHR)

2015: CAT Hong Kong, Beyond the Boundary - Knowing and Concerns Intersex (BBKCI)

2015: CAT Austria, Verein Intersexueller Menschen Österreich VIMÖ, Zwischengeschlecht.org

2015: CAT Denmark, Ditte Dyreborg, Zwischengeschlecht.org

2016: CRC Ireland, Gavan Coleman, Zwischengeschlecht.org

2016: CRC France, Vincent Guillot, Zwischengeschlecht.org
D. What is Intersex?

1. Variations of Sex Anatomy

Intersex persons, in the vernacular also known as hermaphrodites, or medically as persons with “Differences[32] of Sex Development (DSD),” are people born with “atypical” sex anatomies and reproductive organs, including

a) “ambiguous genitalia”, e.g. “enlarged” clitoris, urethral opening not on the tip of the penis, but somewhere below on the underside of the penis (Hypospadias), fused labia, absence of vagina (vaginal agenesis, or Mayer-Rokitansky-Küster-Hauser syndrome MRKH), unusually small penis or micropenis, breast development in “males”; and/or

b) atypical hormone producing organs, or atypical hormonal response, e.g. a mix of ovarian and testicular tissue in gonads (ovotestes, “True Hermaphroditism”), the adrenal gland of the kidneys (partly) producing androgens (e.g. testosterone) instead of cortisol (Congenital Adrenal Hyperplasia CAH), low response to testosterone (Androgen Insensitivity Syndrome AIS), undescended testes (e.g. in Complete Androgen Insensitivity Syndrome CAIS), little active testosterone producing Leydig cells in testes (Leydig Cell Hypoplasia), undifferentiated streak gonads (Gonadal Dysgenesis GD if both gonads are affected, or Mixed Gonadal Dysgenesis MGD with only one streak gonad); and/or

c) atypical genetic make-up, e.g. XXY (Klinefelter Syndrome), X0 (Ullrich Turner Syndrome), different karyotypes in different cells of the same body (mosaicism and chimera).

Variations of sex anatomy include

- “atypical characteristics” either on one or on more of the above three planes a)–c),
- or, while individual planes appear “perfectly normal”, together they “don’t match”, e.g. a newborn with male exterior genitals but an uterus, ovaries and karyotype XX (some cases of Congenital Adrenal Hyperplasia CAH), or with female exterior genitals but (abdominal) testicles and karyotype XY (Complete Androgen Insensitivity Syndrome CAIS).

While many intersex forms are usually detected at birth or earlier during prenatal testing, others may only become apparent at puberty or later in life.

Everybody started out as a hermaphrodite: Until the 7th week of gestation, every fetus has “indeterminate” genitals, two sets of basic reproductive duct structures, and bipotential gonads. Only after the 7th week of gestation, fetuses undergo sexual differentiation mostly resulting in typically male or female sex anatomy and reproductive organs (see Figure 1). However, with some fetuses, sex development happens along a less common pathway, e.g. due to unusual level of certain hormones, or an unusually high or low ability to respond to them, resulting in intersex children born with in-between genitals (see Figure 2) and/or other variations of sex anatomy.

For more information and references on genital development and appearance, please see 2014 CRC NGO Report (A 2–3, p. 8–10.).[33]
Figure 1 “Genital Development Before Birth”

Figure 2 “Genital Variation” (Diagrams 1–6 corresponding to Prader Scale V–0)
2. How common is Intersex?

Since hospitals, government agencies and health insurances covering intersex surgeries on children until the age of 20, refuse to disclose statistics and costs, there are no exact figures or statistics available. Also, the definition of intersex is often arbitrarily changed by doctors and government agencies in order to get favourable (i.e. lower) figures. Ultimately, all available numbers are mere estimates and extrapolations. Intersex persons and their organisations have been calling for independent data collection and monitoring for some time, however to no avail.

An often quoted number is 1:2000 newborns, however this obviously disregards variations of sex anatomy at risk of “masculinising corrections” (hypospadias). In medical literature, often two different sets of numbers and definitions are given depending on the objective:

a) **1:1000** if it’s about getting access to new patients for paediatric genital surgery, and

b) **1:4500 or less** if it’s about countering public concerns regarding human rights violations, often only focusing on “severe cases” while refusing to give total numbers. On the other hand, researchers with an interest in criticising the gender binary often give numbers of up to “as high as 2%”.

However, from a human rights perspective, the crucial question remains: How many children are at risk of human rights violations, e.g. by non-consensual, medically unnecessary, irreversible, cosmetic genital surgeries or other similar treatments justified by a psychosocial indication? Here, the best known relevant number is **1:500 – 1:1000 children are submitted to (often repeated) non-consensual “genital corrections”**.

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35 Rainer Finke, Sven-Olaf Höhne (eds.) (2008), Intersexualität bei Kindern, Preface, at 4

36 e.g. “fewer than 2 out of every 10,000 births”, Leonard Sax (2002), How common is intersex? a response to Anne Fausto-Sterling, The Journal of Sex Research 39(3):174-178, at 178


38 Intersex Society of North America (ISNA), How common is intersex?, [http://www.isna.org/faq/frequency](http://www.isna.org/faq/frequency)
3. Intersex is NOT THE SAME as LGBT

Unfortunately, there are several harmful misconceptions about intersex still prevailing in public, some of which are LGBT-related, e.g. if intersex, and/or intersex status, are represented as a sexual orientation (like gay or lesbian), and/or as a gender identity, as a subset of transgender, as the same as transsexuality,\(^39\) or as a strange, peculiar form of sexual preferences.

The underlying reasons for such misconceptions include lack of public awareness of the situation of real-life intersex persons and the real-life problems they’re facing, as well as – often despite best intentions – a long history of (political) appropriation of intersex going back to the 19th century, including leading LGBT proponents, scholarly authorities and/or interest groups instrumentalising intersex as a means to an end for their own agenda, and/or presenting themselves as intersex and speaking publicly for intersex people.

While some intersex persons position themselves within an LGBT context and many intersex organisations collaborate with LGBT groups on an equal footing to address e.g. discrimination issues, intersex persons and their organisations, as well as their allies, again and again have spoken out clearly against instrumentalising intersex issues as a means for other ends,\(^40\) maintaining that intersex stands for distinct and unique physical variations, and intersex status is not about gender identity or sexual orientation.

\(^{39}\) E.g. the Swiss Federal Government in 2011 in answers to parliamentary questions consistently described intersex as “True and Untrue Transsexualism”, e.g. 11.3286, [http://www.parlament.ch/d/suche/seiten/geschaefte.aspx?gesch_id=20113286](http://www.parlament.ch/d/suche/seiten/geschaefte.aspx?gesch_id=20113286)

• Gabriele Dietze (2006), Schnittpunkte. Gender Studies und Hermaphroditismus, in: Gabriele Dietze, Sabine Hark (eds.): Gender kontrovers. Genealogie und Grenzen einer Kategorie, 46-68, at 56
Although intersex children born with variations of sex anatomy may face several problems, in the “developed world” the most pressing are the ongoing Intersex Genital Mutilations, which present a distinct and unique issue constituting significant human rights violations, which are different from those faced by the LGBT community. Therefore human rights violations of intersex people cannot be addressed properly by framing and addressing them as LGBT issues, but need to be adequately addressed in a separate section as specific intersex issues.

4. Terminology

There is no terminology universally accepted by all persons concerned. All current terms were or are used by medicine in connection with non-consensual, medically not necessary “genital corrections” (see Annexe “Historical Overview”), and/or as insult or verbal abuse in society, and/or have other negative connotations – but all have also been (re-)claimed by persons concerned and their organisations:

**Intersex**
+ Term most frequently used by persons concerned, especially human rights related; reclaimed since 1993.
− In public often leads to misconceptions like “intersex is a sexual orientation,” “intersexuality is a sexual preference”, etc.; “Intersexual Constitution” was a racist/nazi medical diagnosis 1920s–1950s, “Intersex Disorders” was the predominant medical term 1960s–2005.

*Please do say: Intersex child, she’s intersexed, this is an intersex human rights issue.*
*Please don’t say: Intersexuality, intersexual, intersexuals – this is disregarded by many persons concerned today and bound to foster misconceptions.*

**Hermaphrodite**
+ Term most frequently used by public, can dispel misconceptions of intersex as a sexual orientation, sexual preference, etc.; reclaimed since 1994.
− Can lead to misconceptions related to the ancient mythic notion of intersex persons “having both sets of genitals and being able to impregnate themselves”; considered as derogatory by some persons concerned; “(Pseudo) Hermaphroditism” was the medical terminology until the 1950s, though it persisted and is sometimes still used by doctors even today.

*Please do say: Intersex persons, in the vernacular also known as hermaphrodites, are people with variations of sex anatomy.*
*Please don’t say: Are you a hermaphrodite? What do your genitals look like?*

**DSD (Disorders of Sex Development vs. Differences of Sex Development)**
“Disorders of Sex Development”, mostly referred to by the acronym “DSD”, is the current medical term, introduced at the “Chicago Consensus Conference 2005” with limited input by persons concerned, but in an intransparent way and without proper consultation. The new nomenclature also included a new taxonomy based on karyotype and focused on conditions (instead of the persisting “Pseudo Hermaphrodite” taxonomy based on gonadal

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status and focused on “male” and “female”), also the new taxonomy was supposed to more clearly include genital variations irrespective of gender of rearing issues, such as hypospadias, Klinefelter, and MRKH, reflecting the new definition “congenital conditions in which development of chromosomal, gonadal, or anatomic sex is atypical.”

Furthermore, in some cases a more cautious approach to early surgery was suggested. While the use of an acronym for medical purposes, the new taxonomy focused on conditions, clearer inclusion of all genital variations, and the instances of calling for more caution regarding early surgeries were welcomed by persons concerned and their organisations, the term “disorders” was unequivocally abhorred and condemned within the community, because it frames the persons concerned as in need of being (surgically) “corrected”, or “fixed”, e.g. to “relieve [...] parental distress.” However, clinicians readily embraced “disorders.” “Variations of Sex Development (VSD)” was proposed as a less stigmatising alternative in 2006, but rejected by medicine arguing the acronym VSD was already taken. Nonetheless, another proposal in 2008 of “Differences of Sex Development” keeping the DSD acronym has been equally refused by doctors.

Please do say: Differences of Sex Development (DSD).
Please don’t say: Disorders of Sex Development (DSD).

Words are important, words can hurt – however, more important than a wrong word is the continuous regard – or disregard – of the human rights and dignity of the children concerned.

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47 “It is generally felt that surgery that is performed for cosmetic reasons in the first year of life relieves parental distress and improves attachment between the child and the parents [48–51]; the systematic evidence for this belief is lacking.” Peter A. Lee, Christopher P. Houk, S. Faisal Ahmed, Ieuan A. Hughes, LWPES/ESPE Consensus Group (2006), Consensus statement on management of intersex disorders, Pediatrics 118:e488-e500, at e491, http://pediatrics.aappublications.org/content/118/2/e488.full.pdf
E. IGM Practices – Non-Consensual, Unnecessary Medical Interventions

1. What are Intersex Genital Mutilations?

IGM Practices include non-consensual,\(^{50}\) medically unnecessary,\(^{51,52}\) irreversible,\(^{53}\) cosmetic\(^{54}\) genital surgeries, and/or other similar medical treatments, including imposition of hormones, performed on children with variations of sex anatomy, without evidence of benefit for the children concerned,\(^{55,56}\) but justified by “psychosocial indications [...] shaped by the clinician’s own values”,\(^{57}\) the latter informed by societal and cultural norms and beliefs,\(^{58,59}\) enabling clinicians to withhold crucial information from both patients and parents,\(^{60,61}\) and to submit healthy intersex children to risky and harmful invasive procedures “simply because their bodies did not fit social norms”.\(^{62}\)

\(^{50}\) The United Nations (2013), A/HRC/22/53, at para 77: “Children who are born with atypical sex characteristics are often subject to [...] involuntary sterilization, involuntary genital normalizing surgery, performed without their informed consent, or that of their parents”, http://www.ohchr.org/Documents/HRBodies/HRCouncil/RegularSession/Session22/A.HRC.22.53_English.pdf


\(^{53}\) “2. The surgery is irreversible. Tissue removed from the clitoris can never be restored; scarring produced by surgery can never be undone.” Intersex Society of North America (ISNA) (1998), ISNAs Amicus Brief to the Constitutional Court of Colombia, http://www.isna.org/node/97

\(^{54}\) “It is generally felt that surgery that is performed for cosmetic reasons in the first year of life relieves parental distress and improves attachment between the child and the parents [48–51]; the systematic evidence for this belief is lacking.” Peter A. Lee, Christopher P. Houk, S. Faisal Ahmed, Ieuan A. Hughes, LWPES/ESPE Consensus Group (2006), Consensus statement on management of intersex disorders, Pediatrics 118:e488-e500, at e491, http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2082839/

\(^{55}\) “The final ethical problem was the near total lack of evidence—indeed, a near total lack of interest in evidence—that the concealment system was producing the good results intended.” Alice Domurat Dreger (2006), Intersex and Human Rights: The Long View, in: Sharon Sytsma (ed.) (2006), Ethics and Intersex: 73-86, at 75


\(^{57}\) ibid., at 18 and 15.


\(^{59}\) “In cases of intersex clinicians were intentionally withholding and misrepresenting critical medical information.” Alice Domurat Dreger (2006), Intersex and Human Rights: The Long View, in: Sharon Sytsma (ed.) (2006), Ethics and Intersex: 73-86, at 75


Genital surgery is not necessary for gender assignment, and atypical genitals are not in themselves a health issue.\textsuperscript{63} There are only very few situations where some surgery is necessary for medical reasons, such as to create an opening for urine to exit the body.\textsuperscript{64,65}

In addition to the usual risks of anaesthesia and surgery in infancy, IGM practices carry a large number of known risks of physical and psychological harm, including loss or impairment of sexual sensation, poorer sexual function, painful scarring, painful intercourse, incontinence, problems with passing urine (e.g. due to urethral stenosis after surgery), increased sexual anxieties, problems with desire, less sexual activity, dissatisfaction with functional and aesthetic results, lifelong trauma and mental suffering, elevated rates of self-harming behaviour and suicidal tendencies comparable to those among women who have experienced physical or (child) sexual abuse, impairment or loss of reproductive capabilities, lifelong dependency on daily doses of artificial hormones.\textsuperscript{66,67}

2. Most Frequent Surgical and Other Harmful Medical Interventions

Due to space limitations, the following paragraphs summarise the most frequent and egregious forms only. The injuries suffered by intersex people have not yet been adequately documented.\textsuperscript{68} For a more comprehensive list and sources, see 2014 CRC.NGO Report, p. 63–76.

a) IGM 3 – Sterilising Procedures: Castration / “Gonadectomy” / Hysterectomy / (Secondary) Sterilisation

“At 2 1/2 months they castrated me, and threw my healthy testicles in the garbage bin.” (CRC Case No. 2)

Intersex children are frequently subjected to treatments that terminate or permanently reduce their reproductive capacity. Contrary to doctor’s claims, it is known that the gonads by themselves are usually healthy and “effective” hormone-producing organs, often with complete spermatogenesis [...] suitable for cryopreservation.”\textsuperscript{69} Nonetheless, many still undergo early removal of viable gonads (e.g. testes, ovaries, ovotestes) or other reproductive organs (e.g. uterus), leaving them with “permanent, irreversible infertility and severe mental suffering”\textsuperscript{70} and lifelong metabolic problems. When unnecessary sterilising procedures

\textsuperscript{64} ibid., at 3
\textsuperscript{67} Heinz-Jürgen Voß (2012), Intersexualität – Intersex. Eine Intervention, at 50–65
\textsuperscript{68} Rare examples of publications documenting and reviewing reports by persons concerned include:
- Katrln Karkazis (2008), Fixing Sex: Intersex, Medical Authority, and Lived Experience
- Kathrin Zehnder (2010), Zwitter beim Namen nennen. Intersexualität zwischen Pathologie, Selbstbestimmung und leiblicher Erfahrung
\textsuperscript{69} K. Czeloth et al., “Function of Uncorrected Cryptorchid Testes”, 25th ESPU 2014, online
are imposed on children e.g. to address a low or hypothetical risk of cancer, the fertility of intersex people is not being valued as highly as that of non-intersex people. Survivors often have to pay themselves for adequate Replacement Hormones.

For almost two decades, persons concerned have protested unnecessary sterilising treatments, and denounced non-factual and psychosocial justifications, e.g. “psychological benefit” to removing “discordant” reproductive structures, demanding access to screening for potential low cancer risks instead of preemptive castrations. Even some doctors have been criticising unnecessary intersex gonadectomies for decades, e.g. endocrinologist G. A. Hauser (the “H” in “MRKH Syndrome”) stated, “The castration of patients without a tumour converts symptomless individuals into invalids suffering from all the unpleasant consequences of castration.”

What’s more, psychosocial justifications often reveal underlying racist preconceptions by clinicians (reminiscent of the racist and eugenic medical views of intersex predominant during the 1920s–1950s, but which obviously persist), namely the infamous premise, “We don’t want to breed mutants.” (see 2014 CRC NGO Report, p. 52, 69)

Nonetheless, and despite recent discussions in medical circles, unnecessary gonadectomies and other sterilising treatments persist internationally in University Children’s Hospitals. Only a while ago, when the Rapporteurs criticised unnecessary gonadectomies, a paediatric surgeon replied: “Well, if a CAIS person is living as female, what do they need their testes for anyway?”

b) IGM 2 – “Feminising Procedures”: Clitoris Amputation/“Reduction”, “Vaginoplasty”, Dilation

“I can still remember, how it once felt differently between my legs.” (2014 CRC Case No. 3)

In 19th Century Western Medicine, clitoris amputations a.k.a. “clitoridectomies” on girls were prevalent as a “cure” for a) masturbation, b) hysteria, and c) “enlarged clitoris.” While amputations motivated by a) and b) were mostly abandoned between 1900 and 1945, amputations of “enlarged clitorises” took a sharp rise after 1950, and in the 1960s became the predominant medical standard for intersex children.

For four decades, doctors again and again claimed early clitoris amputation on intersex children would not interfere with orgasmic function. Only in the 1980s–1990s, intersex clitoris amputations were eventually replaced by “more modern” techniques a.k.a. “clitoral reduction” (p. 60), again claimed to preserve orgasmic function, despite persons concerned reporting loss of sexual sensitivity, and/or painful scars – complaints also corroborated by recent medical studies. Tellingly, a current paediatric surgeon’s joke on the topic of potential loss of sexual sensation goes, “They won’t know what they’re missing!”

Despite that in infants there’s no medical (or other) need for surgically creating a vagina “big enough for normal penetration” (“vaginoplasty”), but significant risks of complications (e.g. painful scarring, vaginal stenosis), this is nonetheless standard practice. What’s more, in order to prevent “shrinking” and stenosis, the “corrected” (neo) vagina has to be forcibly dilated

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73 see 2014 CRC NGO Report, p. 57–58
75 Personal communication by a doctor attending the 23rd Annual Meeting of ESPU, Zurich 2012
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by continuously inserting solid objects, a practice experienced as a form of rape and child sexual abuse by persons concerned, and their parents.

Clitoris amputations justified by psychosocial indications were taught in Medical Universities as a suitable “therapy” for intersex children diagnosed with “hypertrophic clitoris” until the 1980s. Despite recent public denials by doctors, hospitals, and health departments, systematic early “clitoris reductions” and “vaginoplasty” performed on intersex infants and justified by psychosocial indications, are still practiced in most University Children’s Clinics throughout the world.

c) IGM 1 – “Masculinising Surgery”: Hypospadias “Repair”

“My operated genital is extremely touch-sensitive and hurts very much when I’m aroused.” (CRC Case No. 1)

Hypospadias is a medical diagnosis describing a penis with the urethral opening (“meatus”, or “pee hole”) not situated at the tip of the penis, but somewhere below on the underside, due to incomplete tubularisation of the urethral folds during prenatal formation of the penis. Hypospadias “repair” aims at “relocating” the urethral opening to the tip of the penis. The penis is sliced open, and an artificial “urethra” is formed out of the foreskin, or skin grafts (p. 59).

Hypospadias per se does not constitute a medical necessity for interventions. The justification for early surgeries is psychosocial, e.g. to allow for “sex-typical manner for urination (i.e. standing for males).” According to a “pilot study”, surgery is “intended to change the anatomy such that the penis looks normal.”

Hypospadias “repair” is notorious for high complication rates of 50% and more, as well as causing serious medical problems where none had been before (e.g. urethral strictures leading to kidney failure requiring dialysis), and frequent “redo-surgeries”. Tellingly, for more than 30 years, surgeons have been officially referring to “hopeless” cases of repeat failed “repair” surgeries as “hypospadias cripples” (i.e. made to a “cripple” by unnecessary surgeries, not by the condition!, p. 59), while in medical publications on hypospadias, “[d]ocumentation on complication rates has declined in the last 10 years” (see 2014 CRC NGO Report, p. 54–56).

For more than 15 years, persons concerned have been criticising impairment or loss of sexual sensitivity. However, doctors still refuse to even consider these claims, let alone promote appropriate, disinterested long-term outcome studies.

Since the “2nd Hypospadias Boom” in the 1990s, hypospadias “repair” is arguably by far the most frequent cosmetic genital surgery done on children with variations of sex anatomy internationally. In University Children’s Hospitals, systematic hypospadias “repair” within the first 18 months of life is still considered common practice for children concerned and raised as boys.


Systematic misinformation, refusal of access to peer support, and directive counselling by doctors frequently prevent parents from learning about options for postponing permanent interventions, which has been criticised by persons concerned and their parents for two decades, seconded by bioethicists, and corroborated by studies, including a recent exploratory study (see 2014 CRC NGO Report, p. 71).


Nonetheless, internationally it’s still paediatricians, endocrinologists and surgeons managing diagnostics and counselling of parents literally from “day one.”78 Parents often complain that they only get access to psychological counselling if they consent to “corrective surgery” first, while doctors openly admit seeking early surgeries to facilitate compliance, e.g. referring to “easier management when the patient is still in diapers” (see 2014 CRC NGO Report, p. 72).

Intersex children are systematically lied to and refused access to peer support in order to keep them in the dark about being born intersex, and, if ever told at all, are sworn to secrecy, e.g. “You are a rarity, will never meet another like yourself and should never talk about it to no one” (see 2014 CRC Report, p. 72), severely compounding shame, isolation and psychological trauma in the aftermath of IGM practices.

e) Other Unnecessary and Harmful Medical Interventions and Treatments

“The assistant called in some colleagues to inspect and to touch my genitals as well.” (CRC Case No. 3)

Other common harmful treatments include (as detailed in the 2014 CRC NGO Report):79

• Forced Mastectomy (p. 70)
• Imposition of Hormones (p. 73)
• Forced Excessive Genital Exams, Medical Display, (Genital) Photography (p. 73)
• Human Experimentation (p. 74)
• Denial of Needed Health Care (p. 75)
• Prenatal “Therapy” (p. 75)
• Selective (Late Term) Abortion (p. 76)
• Preimplantation Genetic Diagnosis (PGD) to Eliminate Intersex Fetuses (p. 76)

3. How Common are Intersex Genital Mutilations?

Same as with intersex births (see above p. 34), (university) hospitals, Government agencies and health insurance covering intersex surgeries on children, refuse to disclose statistics and costs, as well as ignoring repeated calls for independent data collection and monitoring (see below p. 47).

What’s more, doctors, government and other institutions involved in IGM practices, if questioned about statistics, are notorious for going to extreme lengths following established patterns of a) disclosing only tiniest fractions of actual treatments, often arbitrarily changing definitions of intersex and variations of sex anatomies in order to justify favourable (i.e. lower) figures, or b) flatly denying any occurrence or knowledge of IGM Practices, while at the same time the same doctors and hospitals, including such under the auspices of said departments, are continuing to publicly promote and perform them. Or, in the rare cases of studies actually “disclosing” numbers, yet another related tactic involves c) manipulation of statistics. For example the world’s largest outcome study on 439 participants, the 2008 “Netzwerk DSD” intersex study, in official publications only gave a misleading overall total figure of “almost 81% of all participants had at least once surgery [...] most of them before entering school.”80

The only published numbers that include a breakdown by age groups available from the “Netzwerk DSD” intersex study with participation of clinics in Germany, Austria and Switzerland stem from a semi-official 2009 presentation. They reveal that, contrary to declarations by doctors as well as cantonal and federal governments, in the most relevant age groups of 4+ years, 87%–91% have been submitted to IGM surgeries at least once, with increasing numbers of repeat surgeries the older the children get (see Figure 3 above – note, how the table conveniently stops at “>2” surgeries, although, especially with “hypospadias repair”, a dozen or more repeat surgeries are not uncommon).

What’s more, although internationally no official statistics are available, internationally the total number of cosmetic genital surgeries performed on intersex children is known to be still rising.


For more than two decades, persons concerned and sympathetic clinicians and academics have tried to reason with the perpetrators, and for 19 years they’ve been lobbying for legal measures, approaching governments as well as national and international ethics and human rights bodies year after year after year, calling for specific legislation to eliminate IGM practices, and criticising the factual impunity of IGM doctors due to statutes of limitations that – both in criminal and civil law – expire long before survivors of early childhood IGM practices would be able to call a court.

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82 e.g. “The UK National Health Services Hospital Episode Statistics in fact shows an increase in the number of operations on the clitoris in under-14s since 2006”, Sarah M. Creighton, Lina Michala, Imran Mushtaq, Michal Yaron (2014), Childhood surgery for ambiguous genitalia: glimpses of practice changes or more of the same?, Psychology & Sexuality 5(1):34-43, at 38

83 e.g. Italy: “Boom in Surgeries on Children with ‘Indeterminate’ Sex, in Rome 50% Increase during the Last 5 Years, 25% Increase on National Level”, according to Aldo Morrone, Director General of the Ospedale San Camillo-Forlanini di Roma, quoted in: “Boom di bimbi con sesso ‘incerto’, a Roma un aumento del 50 per cento”, leggo.it 20.06.2013, https://web.archive.org/web/20131110133723/http://www.leggo.it/NEWS/ITALIA/boom_di_bimbi_con_sesso_quot_incerto_quot_a_roma_aumentano_del_50_per_centro/notizie/294638.shtml
In 2011, the Committee against Torture (CAT) was the first UN body to recognise the lack of adequate laws ensuring redress and investigations, explicitly calling on Germany to 

“Undertake investigation of incidents of surgical and other medical treatment of intersex people without effective consent and adopt legal provisions in order to provide redress to the victims of such treatment, including adequate compensation.”

In the meantime, the Committee against Torture issued similar recommendations to Switzerland, Austria, Denmark and Hong Kong, repeating its the call for “legislative measures to ensure redress” while adding, “Take the necessary legislative, administrative and other measures to guarantee respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected during infancy or childhood to non-urgent medical or surgical procedures”.

In 2012, the Swiss National Advisory Commission on Biomedical Ethics (NEK-CNE) was the first national body to eventually support the call of survivors for legal measures, in Recommendation 12 explicitly urging a legal review of both criminal law and civil liability implications, as well as for a review of associated statutes of limitations, with explicit reference to Art. 124 Criminal Code (FGM).

Paediatric Surgeon Blaise Meyrat, one of only a handful of paediatric surgeons worldwide refusing to do unnecessary surgeries on intersex children, in 2013 was the first doctor to go on record and frankly admit that in the end only legislation will succeed in ending IGM practices, “It’s a pity that, because of a lack of ethical clarity in the medical profession, we have to get legislators involved, but in my opinion it’s the only solution.” And in 2015, on occasion of the 55th Session of the Committee against Torture, Meyrat added, “Only the fear of the judge will make things change. We need statutes of limitation long enough so that victims may sue as adults.”

In 2013, the survivors’ call for legislative measures was seconded by the Special Rapporteur on Torture (SRT), who in his report on “abuses in health-care settings that may cross a threshold of mistreatment that is tantamount to torture or cruel, inhuman or degrading treatment” explicitly stated:

“Children who are born with atypical sex characteristics are often subject to irreversible sex assignment, involuntary sterilization, involuntary genital normalizing surgery, performed without their informed consent, or that of their parents, “in an attempt to fix their sex”, leaving them with permanent, irreversible infertility and causing severe mental suffering.

These procedures [genital-normalizing surgeries] are rarely medically necessary, can cause scarring, loss


85 here: CAT/C/CHE/CO/7, 14 August 2015, para 20: http://intersex.shadowreport.org/public/CAT_C_CHE_CO_7-Concl-Obs-Switzerland-2015_G1520151.pdf, see also:


of sexual sensation, pain, incontinence and lifelong depression and have also been criticized as being unscientific, potentially harmful and contributing to stigma (A/HRC/14/20, para. 23).”

Also in 2013, this call was again seconded by the Council of Europe (COE) in their Resolution 1952 (2013) “Children’s right to physical integrity”, urging states to “ensure that no-one is subjected to unnecessary medical or surgical treatment that is cosmetic rather than vital for health during infancy or childhood, guarantee bodily integrity, autonomy and self-determination to persons concerned, and provide families with intersex children with adequate counselling and support.”

In 2014, an Interagency Statement on Forced Sterilisation by the WHO and 6 more UN bodies explicitly also criticised IGM practices in general: “Children who are born with atypical sex characteristics are often subjected to cosmetic and other non-medically indicated surgeries performed on their reproductive organs, without their informed consent or that of their parents, and without taking into consideration the views of the children involved.”

In addition, the WHO interagency statement explicitly called for “Remedies and redress”, as well as for “Monitoring and Compliance.”

In 2015, also the Committee on the Rights of the Child (CRC) criticised Switzerland for allowing IGM practices to continue, explicitly highlighting “the lack of remedies and compensation in such cases,” and classifying IGM practices as “violence against children” and as a “harmful practice”, thus clearly implicating the urgent need for legislative measures to eliminate them, namely “to ensure that no-one is subjected to unnecessary medical or surgical treatment during infancy or childhood, guarantee bodily integrity, autonomy and self-determination to children concerned”.

In the same year 2015, in their recommendations to Chile, the Committee on the Rights of the Child added, “ensuring effective remedy for victims, including redress and compensation.” And in 2016, CRC reiterated this criticism of IGM practices in their Concluding Observations for Ireland and France.

In 2015, also the Committee on the Rights of Persons with Disabilities (CRPD), referring to the 2011 CAT Concluding Observations, criticised the failure of “upholding bodily integrity of intersex children”, and urged Germany to “take the necessary measures, including of a legislative nature to [...] implement all the recommendations of CAT/C/DEU/CO/5, para. 20 relevant to intersex children.”

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95 CRPD/C/DEU/CO/1, p. 6–7, paras 37-38: http://tbinternet.ohchr.org/_layouts/treatybody-
Again in 2015, the WHO Report “Sexual health, human rights and the law” reiterated: “It has also been recommended [by human rights bodies and ethical and health professional organizations] that investigation should be undertaken into incidents of surgical and other medical treatment of intersex people without informed consent and that legal provisions should be adopted in order to provide remedies and redress to the victims of such treatment, including adequate compensation.”

Nonetheless, globally so far without even a single exception, states refuse to take legislative action to ensure access to redress for IGM survivors.

5. Lack of Impartial Investigation, Disinterested Review and Research (Art. 12, 13)

Persons concerned and their organisations have stressed for almost two decades “the unreliability of research conducted in the setting where the harm was done”, and stressed the imminent need for impartial, disinterested investigation and research, as called for in Art. 12 CAT and the Committee’s own 2011 Concluding Observations, as well as by the 2012 Swiss National Advisory Commission on Biomedical Ethics (NEK-CNE) (Recommendation 9), the 2013 COE Resolution 1952 (para 7), and the 2014 WHO Interagency Statement.

However, to this day, despite repeated calls for impartial investigation and disinterested research, internationally the only “investigations” taking place are the “research” facilitated by the perpetrators themselves, relying on massive state funding.

The only exception proving the rule is an exceptional preliminary research study “Historic Evaluation of Treatment of Persons with Differences of Sex Development” examining 22 cases of clitoris amputations at the Zurich University Children’s Clinic between 1913 and 1968. This preliminary study was initiated and paid for by the University Children’s Clinic after considerable pressure by intersex NGOs and self-help groups. However, the clinic is still struggling with funding to adequately continue this ground-breaking project constituting a global first, and so far no state body considered supporting it.

On the other hand, currently the European Union and affiliated states are spending millions on exculpating “intersex research projects” facilitated by, and in control of, the perpetrators. “DSD-Life” (see Figure 4, opposite p. 47) and “DSDnet”, two current examples, are conducted by the perpetrators themselves, e.g. in “DSDnet” paediatric endocrinologists and in “DSD-Life” paediatric endocrinologists and paediatric surgeons taking the lead – exactly the professional groups responsible for IGM practices in the first place. If other disciplines are included at all in the “multidisciplinary teams,” like e.g. psychology or bioethics, let alone persons concerned, they only play a secondary role, and are only included at a later stage, and especially persons concerned serve mostly to recruit participants – same as in the precursor projects “Netzwerk DSD” and “EuroDSD”.

external/Download.aspx?symbolno=CRPD/C/DEU/CO/1
96 p. 27, http://apps.who.int/iris/bitstream/10665/175556/1/9789241564984_eng.pdf?ua=1
100 http://www.cost.eu/about_cost/who/%28type%29/5/%28wid%29/1438
101 http://www.cost.eu/domains_actions/bmbs/Actions/BM1303?management
102 http://www.dsdlife.eu/the-group/consortium/, for a more accessible graphic overview of the consortium see: http://stop.genitalmutilation.org/post/IGM-Primer-2-The-Global-Cartel

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What’s more, all of these perpetrator’s “research projects” continue to openly advocate IGM, as well as to promote the usual psychosocial and non-factual justifications, e.g. “DSDnet”:

“Children with DSD may be born with genitalia that range from being atypical to truly ambiguous and the sex assignment process may be extremely challenging for families and health care professionals. Often, multiple surgical interventions are performed for genital reconstruction to a male or female appearance. The gonads are often removed to avoid malignant development.”

On the other hand, to this day an impartial investigation into past and current IGM practices isn’t even considered by any state.

6. Lack of Independent Data Collection and Monitoring (Art. 12, 13)

With no statistics available on intersex births, let alone on surgeries and costs, and perpetrators, governments and health departments colluding to keep it that way as long as anyhow possible, persons concerned as well as civil society lack possibilities to effectively highlight and monitor the ongoing mutilations. What’s more, after realising how intersex genital surgeries are increasingly in the focus of public scrutiny and debate, perpetrators of IGM practices respond by suppressing complication rates, as well as refusing to talk to journalists “on record”.

104 Personal communication by journalist SRF (Swiss National Radio and TV), 2013
F. The Treatment of Intersex Persons as a Violation of the Convention against Torture

“Genital mutilation of intersex children damages genital sensitivity in irreversible ways; it causes postsurgical trauma, and the internalization of brutal prejudices denying or stigmatizing the diversity that in reality human bodies show. [...] The difference in genitalia cannot justify, under any pretext whatsoever, ethical and political hierarchies: cannot justify mutilation, because it never normalizes but does the opposite. For us, mutilation creates a permanent status of human rights violation and inhumanity.”

Mauro Cabral, CESCR NGO Statement 2004

For 23 years now, intersex people from all over the world, and their organisations have been publicly denouncing IGM Practices as destructive of sexual sensation, and as a violation of basic human rights, notably the right to physical integrity. For 19 years, they have lobbied for legislation against IGM Practices to end the impunity of perpetrators due to statutes of limitation. For 18 years, they have been invoking UN Conventions, and for 12 years they have been reporting IGM Practices to the UN as a human rights violation.

In every intersex community, meanwhile several generations of intersex persons, their partners and families, as well as NGOs and other human rights and bioethics experts, have again and again described IGM Practices as a human rights issue, as harmful and traumatising, as torture, as a western form of genital mutilation, as child sexual abuse, and have called for legislation to end it.

The UN Committees CAT, CRC, CRPD, CEDAW, the UN Special Rapporteur on Torture (SRT), the UN Special Rapporteur on Health (SRH), the UN High Commissioner for Human Rights (UNHCHR), the World Health Organisation (WHO), the Council of Europe (COE), and last but not least the Swiss National Advisory Commission on Biomedical Ethics (NEK-CNE) have all recognised the treatment of intersex children as a serious human rights violation, have called for legislative measures (CAT, SRT, NEK-CNE)
1. State Parties’ Commitment to the Prevention of Torture and Cruel, Inhuman or Degrading Treatment (CIDT)

By ratifying the Convention against Torture (CAT), the state parties committed themselves to ensuring that no child within its jurisdiction is subject to torture and other cruel, inhuman or degrading treatment or punishment (CIDT). In addition, state parties may have ratified the Convention on the Rights of the Child (CRC), and the European Convention on Human Rights (ECHR), which both prohibit CIDT, as well as the International Covenant on Civil and Political Rights (ICCPR) which in its Art. 7 contains a similar clause and explicitly includes freedom from forced medical experimentation. The prohibition of torture is absolute and non-derogable.\(^{116}\) All of these Conventions are enforceable statutory law by virtue of their ratification. In addition, many state’s constitutions also ensure the right to life and personal freedom, particularly the right to physical and mental integrity, often explicitly prohibit torture or CIDT, ensure the right of special protection of the integrity of children and young people, as well as equality and non-discrimination.

2. The Treatment of Intersex Persons as Torture

In Article 1 of CAT, torture is defined as:

“any act by which severe pain or suffering, whether physical or mental, is intentionally inflicted on a person for such purposes as obtaining from him or a third person information or a confession, punishing him for an act he or a third person has committed or is suspected of having committed, or intimidating or coercing him or a third person, or for any reason based on discrimination of any kind, when such pain or suffering is inflicted by or at the instigation of or with the consent or acquiescence of a public official or other person acting in an official capacity. It does not include pain or suffering arising only from, inherent in or incidental to lawful sanctions.”

Although many cases of torture happen in detention, torture is no longer understood to constitute solely interrogation, punishment or intimidation of a captive.\(^{117}\) Rather, the definition includes any setting. The UN Special Rapporteur on Torture pointed out in 2008:

“Whereas a fully justified medical treatment may lead to severe pain or suffering, medical treatments of an intrusive and irreversible nature, when they lack a therapeutic purpose, or aim at correcting or alleviating a disability, may constitute torture and ill-treatment if enforced or administered without the free and informed consent of the person concerned.”\(^ {118}\)

In light of this definition, medically unnecessary genital “correction” surgeries and hormone treatments that were not legally consented to by the patient constitute torture in violation of Article 1(1) of the Convention. That is, that such surgeries constitute acts that cause severe pain or suffering (a), they are intentional (b), they serve a specific purpose (c), there is a sufficient nexus with a public official (d) and they are not lawfully sanctioned (e).

\(^{116}\) Art. 2(2) CAT; Nowak/McArthur (2008), Convention Against Torture, Art. 3 para. 200; CAT, General Comment No. 2, CAT/C/GC/2, para. 2-6.

\(^{117}\) Sifris (2010), Conceptualising involuntary sterilisation as “severe pain or suffering” for the purposes of torture discourse, Neth. Qu. HR 28(4), 523-547, at 526.

\(^{118}\) Interim report of the Special Rapporteur on the question of torture and other cruel, inhuman or degrading treatment or punishment, A/63/175, of 28 July 2008, para. 47.
a) Infliction of Severe Pain or Suffering

The infliction of severe pain or suffering on a person can be physical or mental. **Mental suffering** has been defined as the infliction of pain through the creation of a state of anguish and stress by means other than bodily assault. 119 Each circumstance of torture needs to be considered individually, in the context and circumstances, and there is no definitive list of what constitutes a tortuous act. 120

The severity of pain and suffering is relative and therefore has to be evaluated in the specific context. Therefore, the severity of an act that might constitute torture needs to be assessed from an objective perspective that looks at each specific situation and each particular victim and his/her vulnerability. 121 Thereby one needs to take into account different factors, such as the duration of the treatment, its physical/mental effects and the sex, age, state of health of the victim. 122 Thus, the UN Special Rapporteur on Torture has pointed out that children are more vulnerable to the effects of torture as they are in the critical stages of physical and psychological development where they may suffer graver consequences than similarly ill-treated adults. 123 The effects of torture/ill-treatment will also differ according to the age of the child, depending on the readiness of mind. Torture inflicted on a child might leave more long-lasting effects than on an adult. 124 As with children undergoing female genital mutilation (FGM), intersex children undergoing IGM Practices at an early age are in a situation of powerlessness, as they are under the complete control of their parents and have no means of resistance. 125

While the surgery performed on intersex persons will normally involve adequate pain management (anaesthesia), IGM Practices have severe effects on the intersex person’s physical and psychological wellbeing which constitute an infliction of severe pain or suffering:

- **IGM 3: Sterilising Procedures** (see above p. 39) leading to “permanent, irreversible infertility [...] causing severe mental suffering” 126, as well as to the termination of natural hormone production, which also causes mental suffering, 127 and which requires life-long hormone substitution, which also results in severe physical suffering. 128 In its General Recommendation No. 19, the CEDAW Committee notes that compulsory sterilisation adversely affects women’s mental health, and likewise will it affect a man’s mental health. 129 Moreover, in a recent case involving the sterilization of a Hungarian Romani woman without her knowledge or informed consent, the Com-

121 Ibid., p. 28.
122 ECHR, Ireland v UK (1978) 2 EHRR 25, para. 162.
128 Ibid., at 18.
mittee noted the profound impact that the sterilization had on her life, resulting in her and her partner being treated medically for depression and psychological trauma. The Special Rapporteur on Torture has also taken up the subject in strong words. The sterilization of women without their consent has been recognized as a breach of the prohibition on torture. Consequently, the Committee against Torture, the Special Rapporteur on Torture and the WHO plus 6 more UN bodies have issued strong statements specifically criticising forced sterilising procedures on intersex persons.

- **IGM 2: “Feminising” Surgical and Other Procedures** (see above p. 40) including removal or recession of the clitoris, vaginal surgery and dilation, leading to impairment or loss of genital sensitivity, painful intercourse, sexual dysfunction and suicidal tendencies, causing severe physical and mental suffering.

The removal or recession of the clitoris has been considered in international law as part of Female Genital Mutilation (FGM). The UN Special Rapporteur on Torture, the UN Special Rapporteur on Violence against Women and the Human Rights Committee have made it clear that FGM constitutes torture and that, from a human rights perspective, the medicalisation of FGM – its performance in clinical surroundings – does not make this practice more acceptable. This also holds for the mutilation of the clitoris of intersex children or adults as part of unnecessary feminising cosmetic surgery which, like FGM, is performed for purely cultural reasons. Accordingly, the Committee on the

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132 CCPR General Comment No. 28 (2000) on article 3 (The equality of right between men and women), para. 20. See also Concluding Observations on Slovakia, CCPR/CO/78/SVK, para. 12; on Japan, CCPR/C/79/ADD.102, para. 31; and on Peru, CCPR/CO/70/PER, para. 21. See also CAT, Concluding Observations on Peru, CAT/C/PER/CO/4, para. 23.
138 This procedure is also called Female Genital Cutting (FGC). The World Health Organization defines FGM as “all procedures that involve partial or total removal of the external female genitalia, or other injury to the female genital organs for non-medical reasons” and classifies it into four types, one of which is clitoridectomy.
139 Report of the Special Rapporteur on Torture, A/HRC/7/3, paras. 53, 54; Report of the Special Rapporteur on Violence against Women, E/CN.4/2002/83, para. 6 (severe pain and suffering element of CAT definition); see also A/HRC/4/34, para. 56. Breach of Art. 7 ICCPR: see CCPR general comment No. 28 (2000) on article 3 (The equality of rights between men and women), para. 11; see also Concluding Observations on Uganda, CCPR/CO/80/UGA, para. 10; Mali, CCPR/CO/77/MLI, para. 11; Sweden, CCPR/CO/74/SWE, para. 8; Yemen, CCPR/CO/84/YEM, para. 11.
Rights of the Child explicitly considered “medically unnecessary surgical and other procedures on intersex children, which often entail irreversible consequences and can cause severe physical and psychological suffering” a “harmful practice”.  

**Genital dilation** is described as a very a painful experience. Other than the above treatments which are performed under anaesthesia, intersex persons are dilated repeatedly to prevent the downsizing of the tissue. The repeated insertion of a solid object into a young person’s vagina does not only pain the aggrieved persons, but it is also highly traumatic. Such invasions of the body, performed without the acquiescence of the victim, constitute rape. The ICTR in its Akayesu judgement, has established that in international law, rape is not limited to the penetration of the vagina with a penis but encompasses other bodily invasions, including with objects or with other parts of the body. The Inter-American Court of Human Rights thus considered a “finger vaginal ‘examination’ [...] sexual rape that due to its effects constituted torture”, an invasion similar to what is endured during dilation. As rape “leaves deep psychological scars on the victims which do not respond to the passage of time as quickly as other forms of physical and mental violence”, it has been found to constitute torture in many international settings. Intersex people who have endured dilation as children often report to reject any kind of penetration at adulthood, and to experience any kind of physicality as torment.

The most severe mental suffering, regardless of what form of surgery was performed, results in suicidal tendencies. In a study conducted in Hamburg, Germany, 50% of those that had been subjected to irreversible surgical interventions were found to contemplate suicide. Another study found the elevated rates of self-harming behaviour and suicidal tendencies among “DSD” individuals comparable to those among women traumatised with physical or sexual abuse.

In addition to the Committee on the Rights of the Child declaring IGM a “harmful prac-

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148 Schützmüller et al. (2009), Psychological distress, suicidal tendencies, and self-harming behaviour in adult persons with different forms of intersexuality, Arch Sex Behav. 2009 Feb;38(1):16-33.
tice” (see above), also the Committee against Torture,\(^{149}\) the Special Rapporteur on Torture\(^ {150}\) and the WHO\(^ {151}\) plus 6 more UN bodies\(^ {152}\) have issued strong statements specifically criticising unnecessary surgical procedures on intersex persons.

- **IGM 1: “Masculinising” Surgical Procedures** (see above p. 41) are regularly resulting in severe complications,\(^ {153}\) obviously leading to impairment or loss of genital sensitivity, painful intercourse, sexual dysfunction and suicidal tendencies, causing severe physical and mental suffering. Also in what doctors refer to as “successful cases”, persons concerned report impairment of sensation.\(^ {154}\) Thus, the criticisms by CRC (“harmful practice”), as well as those by CAT, SRT, WHO and 6 more UN bodies referenced above under “Feminising” Surgical and Other Procedures” also apply to “masculinising” procedures accordingly.

b) Intention

The Special Rapporteur on Torture points out that intent can be implied where the act had a specific purpose,\(^ {155}\) namely where a person has been discriminated against on the basis of disability.\(^ {156}\) Intent and purpose do not require a subjective inquiry into the motivation of the perpetrators, but rather an objective determination under the circumstances.\(^ {157}\) The Rapporteur emphasises this in the context of medical treatment, where such discriminations are often “masked as ‘good intentions’ on the part of health professionals”.\(^ {158}\) Where individuals are discriminated against on the basis of bodily features pathologised as “disorders of sex development” (DSD) in medical terms, this discrimination will thus imply intent.


\(^{156}\) Interim report of the Special Rapporteur on the Torture, UN Doc. A/63/175, para. 30.

\(^{157}\) Ibid. para. 49.

\(^{158}\) CAT, General Comment No. 2 (2007), CA/C/GC/2, para. 9.
Clearly, surgery on intersex persons is **always intentionally performed** and not merely the result of negligence. Doctors are also aware that there is usually no medical indication for such surgery but nonetheless approve of the irreversibility of the treatments and the heavy consequential physical and psychological damages of their patients. The physical and mental suffering caused by IGM Practices is well-established in medical literature (see above a). It is thus **foreseeable** to those intentionally inflicting the treatment that severe pain and suffering will ensue.

It does not detract from the intention that doctors perform surgery for **well-meant purposes**. This has been established in a case where a medical team discriminated against a person with disabilities. The same is true for intersex persons where doctors believe to prevent cancer or social ostracism. The fact that there is no medical justification for the ill-treatment means that good intentions cannot prevent the treatment from constituting torture.

c) **Purpose of Discrimination**

Article 1 of CAT requires that the pain or suffering be inflicted for one of the enumerated purposes, i.e. for the extraction of information or confession, punishment, intimidation and coercion, “or for any reason based on discrimination of any kind”.

The Committee against Torture emphasised that the protection of certain minority or marginalised individuals or populations especially at risk of torture is part of the State obligation to prevent torture. State parties must make sure that with respect to the Convention, their laws are in practice applied to all persons, “regardless of [...] gender, sexual orientation, transgender identity, mental or other disability, health status, [...]”. This includes fully prosecuting and punishing all acts of violence and abuse against these individuals and implementing positive prevention and protection measures.

On the basis of their “indeterminate sex,” intersex children are singled out for experimental harmful treatments, including surgical “genital corrections” and sterilising procedures, that would be “considered inhumane” on “normal” children, by reverting to a “monster approach” implying intersex children are “so grotesque, so pathetic, any medical procedure aimed at normalizing them would be morally justified”, so that, according to a specialised surgeon, “any cutting, no matter how incompetently executed, is a kindness.”

By means of surgery, intersex children are penalised compared to “normal” infants, even where the perpetrator has benign intentions.

d) **Involvement of a State Official**

As underlined by the Committee, the prohibition of torture must be enforced in all institutions, including hospitals that engage in the care of children. The Special Rapporteur on Torture underlined that the obligation to prevent torture extends “to doctors, health professionals

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159 Ibid.
160 CAT, General Comment No. 2 (2008), CAT/C/GC/2, para. 21.
164 CAT, General Comment No. 2 (2008), CAT/C/GC/2, para. 15.
and social workers, including those working in private hospitals [or] other institutions.”

The medical ill-treatment of intersex persons is attributable to state as it is committed by or at the instigation of or with the acquiescence of a person acting in an official capacity, either by way of involvement of public hospitals, universities and insurances, or by the failure of the State to exercise due diligence to protect this group of citizens from torture.

e) Lawful Sanction

Surgery performed on an intersex child or adult does not constitute a sanction. It is therefore not covered by the exception clause.

3. The Treatment of Intersex Persons as Cruel, Inhuman or Degrading Treatment (CIDT)

Article 16 of the Convention commits each State Party to the prevention of:

“other acts of cruel, inhuman or degrading treatment or punishment which do not amount to torture as defined in article 1, when such acts are committed by or at the instigation of or with the consent or acquiescence of a public official or other person acting in an official capacity.”

Acts which fall short of torture are thus still prohibited if they amount to cruel, inhuman or degrading treatment or punishment. This is the case if the treatment does not reach the requisite threshold of severity, or if the suffering was inflicted negligently (see above 2.b).

Thus, if it is considered that the treatment that intersex persons suffer does not meet the severity threshold of Article 1 of the Convention, it certainly meets the threshold of Article 16. If it is considered that this suffering is not foreseeable to the surgeons, the insurance companies or the State, this lack of consideration constitutes negligence sufficient for Art. 16. As to State involvement and due diligence, the same applies as above. A discriminatory or other purpose is not required for CIDT.

Thus, even if it is considered that the treatment of intersex people does not constitute torture, it certainly constitutes cruel, inhuman and degrading treatment which is equally prohibited by the Convention in absolute and non-derogable terms. According to the Committee’s General Comment 3, for CIDT also Article 14 applies.

4. Lack of Legislative Provisions to Ensure Protection from IGM Practices, Impunity of the Perpetrators (Art. 2, 14, 16)

Art. 2 of the Convention obliges State parties to “take effective legislative, administrative, judicial or other measures to prevent acts of torture in any territory under its jurisdiction.” General Comment 2 states, “The obligation to prevent ill-treatment in practice overlaps with and is largely congruent with the obligation to prevent torture,” and similarly obliges State parties to “to eliminate any legal or other obstacles that impede the eradication of torture and ill-treatment; and to take positive effective measures to ensure that such conduct and any recurrences thereof are ef-

165 Interim report of the Special Rapporteur on Torture, UN Doc. A/63/175, para. 51, referencing CAT General Comment No. 2 (2008), para. 17. See also A/HRC/7/3, para. 31.

See also: Report of the Special Rapporteur on Torture, A/HRC/22/53, 1 February 2013, paras 17, 18, 20, 23, 24, 32, 38


167 Committee against Torture (2012), General comment No. 3, CAT/C/GC/3, para. 1.
Accordingly, with regards to IGM practices, the Committee already explicitly recognised the obligation for State parties to “Take the necessary legislative, administrative and other measures to guarantee respect for the physical integrity and autonomy of intersex persons and to ensure that no one is subjected during infancy or childhood to non-urgent medical or surgical procedures.”

However, to this day Malta is still the only nation to at least formally outlaw IGM practices— but only in Civil Law, and without addressing accountability, or immunity of the perpetrators, nor sanctions, nor IGM performed abroad (arguably the majority of Maltese intersex children are sent to Italy for IGM treatments), nor access to redress and justice for victims. Thus, globally the lack of appropriate legislative measures to effectively eliminate IGM practices prevails, as well as the factual impunity of the perpetrators.

This situation is not in line with state parties’ obligations under the Convention.

5. Obstacles to Redress, Fair and Adequate Compensation

Articles 12 and 13 of the Convention require that the State provide the means for an impartial inquiry into allegations of torture or CIDT (Art. 16 CAT). Article 14 requires an enforceable right to redress, fair and adequate compensation, including the means for as full rehabilitation as possible. However, intersex people encounter serious difficulties pursuing their rights.

The statutes of limitation prohibit survivors of early childhood IGM Practices to call a court long before they become adults, despite the fact that persons concerned often do not find out about their medical history until much later in life, and severe trauma caused by IGM Practices often prohibits them to act in time once they do. Globally, states refuse to take legislative action to change that, and refuses to initiate impartial investigations, as well as data collection, monitoring, and disinterested research. In addition, hospitals are often unwilling to provide access to patient’s files.

This situation is not in line with state parties’ obligations under Articles 12–14 of the Convention.

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168 Committee against Torture (2008), General comment No. 2, CAT/C/GC/2, para. 3-4.
171 Globally, no survivor of early surgeries ever managed to have their case heard in court. All relevant court cases (3 in Germany, 1 in the USA) were either about surgery of adults, or initiated by foster parents.
6. Conclusion: Internationally, States are Failing their Obligations towards Intersex People under the Convention against Torture

The surgeries and other harmful treatments intersex people endure cause severe physical and mental pain. Doctors perform the surgery for the discriminatory purpose of making a child fit into societal and cultural norms and beliefs, although there is plenty of evidence on the suffering this causes. State parties are responsible for these violations amounting to torture or at least ill-treatment, committed by often publicly funded doctors, clinics, and universities, as well as in private clinics, all relying on money from often mandatory health insurance, and public grants. Although in the meantime the pervasiveness IGM practices is common knowledge, and most state parties will have been repeatedly called to action both on state, federal, and international level, nonetheless they fail to prevent these grave violations both in public and in private settings, but allow the human rights violations on intersex children and adolescents to continue unhindered and the perpetrators continue with impunity, often condoning and justifying the practice against criticism by human rights bodies and intersex human rights defenders. Nowhere, victims of IGM practices are allowed access to redress and justice.

Such state parties are thus in breach of their obligation to take effective legislative, administrative, judicial or other measures to prevent acts of torture (Art. 2 CAT) or other forms of cruel, inhuman or degrading treatment (Art. 16 CAT, GC 2).

Even where torture is a punishable offense in state law (Art. 4 CAT), victims IGM practices encounter severe obstacles in the pursuit of their right to an impartial investigation (Art. 12, 13 CAT), and to redress, fair and adequate compensation, including the means for as full rehabilitation as possible (Art. 14 CAT).

Globally, state parties’ efforts on education and information regarding the prohibition against torture in the training of medical personnel are grossly insufficient with respect to the treatment of intersex people (Art. 10 CAT).
"Hypospadias," i.e. when the urethral opening is not on the tip of the penis, but somewhere on the underside between the tip and the scrotum, is arguably the most prevalent diagnosis for cosmetic genital surgeries. Procedures include dissection of the penis to "relocate" the urinary meatus. Very high complication rates, as well as repeated "redo procedures" — "5.8 operations (mean) along their lives ... and still most of them are not satisfied with results!"

Nonetheless, clinicians recommend these surgeries without medical need explicitly "for psychological and aesthetic reasons." Most hospitals advise early surgeries, usually "between 12 and 24 months of age." While survivors criticise a.o. impairment or total loss of sexual sensation and painful scars, doctors still fail to provide evidence of benefit for the recipients of the surgeries.

**Hypospadias Surgery**

- **Elbakry (BJUI 88: 590-595, 2001): 42% complications**
  - 5 breakdowns (7%)
  - 17 fistulae (23%)
  - Urethral strictures (9%)
  - Urethral diverticulae (4%)

- **Asopa / Duckett tube**
  - 3.7% (El-Kasaby J Urol 136: 643-644, 1986)
  - 69% (Parsons BJU 25: 186-188, 1984)
  - 15% (Duckett - 1986)

**Hypospadias - Procedures for cripple hypospadias**

- No standardized procedures
- Personal experience of the surgeon
- Importance of a uro-endocrine approach of complex cases to increase the healing abilities of the penile tissues

**Onlay island flap urethroplasty**

- Rectangular skin incision around the fistula orifice, often lateral
- Dissection and excision of the fistula tract
- Urethral suture
- Multilayer cover with well-vascularized tissue (tunica vaginalis, darts, dorsal subcutaneous flap ...)
- Problem: coronal fistula
  - +++; Prefer redo urethroplasty
- Suprapubic diversion?

**Treatment of isolated fistulae**

- **Elbakry**

**Onlay / Duckett - results**

- **Official Diagnosis “Hypospadias Cripple”**
  = made a "cripple" by repeat cosmetic surgeries

**Hypospadias - Conclusions**

- Hypospadias surgery remains a surgical challenge
- Long-term results are poorly reported
- Essential joint uro-endocrine approach
- Psychological consequences poorly assessed
- Informing parents is crucial: 50% of all hypospadias will require further surgical attention during their life.
- Research: Essential role of the placenta / Penile growth factors / healing factors / blood supply ...

**IGM 2 – “Feminising Surgery”: “Clitoral Reduction”, “Vaginoplasty”**

Partial amputation of clitoris, often in combination with surgically widening the vagina followed by painful dilation. “46,XX Congenital Adrenal Hyperplasia (CAH)” is arguably the second most prevalent diagnosis for cosmetic genital surgeries, and the most common for this type (further diagnoses include “46,XY Partial Androgen Insufficiency Syndrome (PAIS)” and “46,XY Leydig Cell Hypoplasia”).

Despite numerous findings of impairment and loss of sexual sensation caused by these cosmetic surgeries, and lacking evidence for benefit for survivors, current guidelines nonetheless advise surgeries “in the first 2 years of life”, most commonly “between 6 and 12 months,” and only 10.5% of surgeons recommend letting the persons concerned decide themselves later.

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**Note Caption 8b:** “Material shortage” [of skin] while reconstructing the praeputium clitoridis and the inner labia.

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**Source (above):** Christian Radmayr: *Molekulare Grundlagen und Diagnostik des Intersex*, 2004

**Source (above):** Finke/Höhne: *Intersexualität bei Kindern*, 2008

Note Caption 8b: “Material shortage” [of skin] while reconstructing the praeputium clitoridis and the inner labia.

IGM 3 – Sterilising Surgery: Castration / “Gonadectomy” / Hysterectomy

Removal of healthy testicles, ovaries, or ovotestes, and other potentially fertile reproductive organs. “46,XY Complete Androgen Insufficiency Syndrome (CAIS)” is arguably the 3rd most common diagnosis for cosmetic genital surgeries, other diagnoses include “46,XY Partial Androgen Insufficiency Syndrome (PAIS)”, male-assigned persons with “46,XX Congenital Adrenal Hyperplasia (CAH)”, and other male assigned persons, who have their healthy ovaries and/or uteruses removed.

Castrations usually take place under the pretext of an allegedly blanket high risk of cancer, despite that an actual high risk which would justify immediate removal is only present in specific cases (see table below), and the admitted true reason is “better manageability.” Contrary to doctors claims, it is known that the gonads by themselves are usually healthy and “effective” hormone-producing organs, often with “complete spermatogenesis [...] suitable for cryopreservation.”

Nonetheless, clinicians still continue to recommend and perform early gonadectomies – despite all the known negative effects of castration, including depression, obesity, serious metabolic and circulatory troubles, osteoporosis, reduction of cognitive abilities, loss of libido. Plus a resulting lifelong dependency on artificial hormones (with adequate hormones often not covered by health insurance, but to be paid by the survivors out of their own purse).

<table>
<thead>
<tr>
<th>Table 1. Prevalence of type II GCT in various forms of DSD</th>
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<tbody>
<tr>
<td><strong>Risk</strong></td>
</tr>
<tr>
<td>High</td>
</tr>
<tr>
<td>46,XY GD</td>
</tr>
<tr>
<td>Frasier syndrome</td>
</tr>
<tr>
<td>Denys-Drash syndrome</td>
</tr>
<tr>
<td>45,X/46,XY GD</td>
</tr>
<tr>
<td>Intermediate</td>
</tr>
<tr>
<td>17β-hydroxysteroid dehydrogenase deficiency</td>
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<tr>
<td>Low</td>
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<tr>
<td>Ovotesticular DSD</td>
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<tr>
<td>Unknown</td>
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</tbody>
</table>

GD = Gonadal dysgenesis; PAIS = partial androgen insensitivity syndrome; CAIS = complete androgen insensitivity syndrome. *
 Might reach more than 30%, if gonadectomy has not been performed.


PAIS

- Bilateral gonadectomy
- Skin Biopsy for genetics study of androgen receptors
- Female gender assignment
- Feminizing genitoplasty performed age 6 months

Buenos Aires 1925: Medical Display, “Trophy Shots”, and Cosmetic Genital Surgeries on Children

“Las deformidades de la sexualidad humana” by Carlos Lagos García (1880-1928) is arguably the first modern medical book dedicated exclusively to “genital abnormalities” and their surgical “cure”. It was highly influential both in Europe and the Americas, pioneering forced medical display, “trophy shots” of amputated healthy genitals and reproductive organs, and advocating cosmetic surgeries on little children, both “feminising” and “masculinising” – expressly without actual medical necessity, but as “correction” for “anomalies”.

Hugh Hampton Young (1870-1945), "The Father of American Urology", also pioneered Intersex Genital Mutilations at the Johns Hopkins University Hospital in Baltimore – a fact nowadays often "neglected" in official hagiographies, despite that Young's disturbing textbook "Genital Abnormalities, Hermaphroditism, and Related Adrenal Diseases" was considered a breakthrough by his colleagues and was received globally. It saw two updated revisions, edited by Young's successors Howard W. Jones and William Wallace Scott, in 1958 and 1971 under the slightly modified title "Hermaphroditism, Genital Anomalies, and Related Endocrine Disorders", and still contained many of Young's original step by step tutorials e.g. of "Plastic operations to construct a vagina and amputate hypertrophied clitoris", or how to otherwise freely "cut up and re-assemble" so called "Genital Abnormalities." Also the Fig. 64 above right showing the tragically mutilated young person "Case 5 / BUI 14127" appeared again in Jones' and Scott's editions, although erroneously attributed to another "Case." For the 1958 edition, Young's colleague at Johns Hopkins and the "inventor" of systematic cosmetic genital surgeries on children, Lawson Wilkins, contributed a foreword, praising Young's original 1937 edition as a "classic."

Paris 1939: “Embarrassing Erections”, yet more “Trophy Shots”, and even younger Children submitted to Cosmetic Genital Surgeries

Louis Ombrédanne (1871-1956) set the standard for “Hypospadias Repairs” a.k.a. “masculinising corrections” for more than 50 years, and even more so for medical musings on allegedly “embarrassing and maybe even painful erections” of “enlarged clitorises” (note how he’s asking himself, NOT his patients), and was a teacher of Swiss paediatric surgeon Max Grob (Zurich University Children’s Hospital). Ombrédanne’s “Hermaphrodites and Surgery” drew heavily on Carlos Lagos García, as well as featuring a “personal observation” by García’s Brother Alberto Lagos García involving a “partial resection of the hypertrophied clitoris” in combination with “continued vaginal dilatations” on a “girl aged three years” (p. 248), and was received internationally from Zurich to Baltimore and beyond.


Geneticist Richard Goldschmidt (1878–1958), before serving as director at the “Kaiser-Wilhelm-Institut für Biologie” in Berlin, coined the terms “Intersex” and “Intersexuality” when internationally publicising his experiments of crossbreeding “different geographic races” of gypsy moths during a stay in the USA (first in English, later in German), claiming to be able to produce “hermaphroditic” a.k.a. “intersex” specimens of any grade and shape at will, and thereafter extrapolating his findings to humans. Of Jewish descent, Goldschmidt was forced to leave the “Kaiser-Wilhelm-Institute” in 1936 and emigrated to the United States. Despite Goldschmidt’s downplaying the “racial” background of his findings since the early 1930’s and later renouncing the underlying genetic theories altogether, the term “Intersex” and its racial implications prevailed. The derived diagnosis “Intersexual Constitution” (published by Austrian Gynaecologist Paul Mathes and Swiss Gynaecologist Hans Guggisberg in 1924), allegedly most frequent amongst “Jews,” and associated with “biological inferiority”, mental illnesses (see above “schizoid”), “hypertrophied clitoris,” and a strict verdict “not fit for marriage,” was particularly popular among prominent eugenicists and Nazi doctors, amongst others Fritz Lenz, Lothar Gottlieb Tirala, Robert Stigler, Wilhelm Weibel, Walther Stoeckel, and kept being used in publications years after World War II.

Baltimore 1950: From Experimentation to Medical Extermination

Lawson Wilkins (1894-1963), "The Father of Pediatric Endocrinology", and teacher of the famous Swiss paediatric endocrinologist Andrea Prader in 1950, was also the “inventor” of systematic cosmetic genital surgeries on children. As his monograph illustrates, in 1950 at Johns Hopkins in Baltimore, any child diagnosed “not normal” was submitted to drastic “Genital Corrections”, either “feminising” or “masculinising”. Often John Money gets erroneously credited as having “invented” the systematic mutilations, however, it was Wilkins (and Prader) who started systematic surgeries; Money “only” delivered a “scientific rationale” five years after the fact.

Zurich 1957: Prader Scales, “Surely Justified” Clitoris Amputations, and even more “Embarassing” Psychosocial Indications

Swiss paediatric surgeon Max Grob (1901-1976), trained in Paris by Ombrédanne, served as director of the Zurich University Children’s Hospital’s paediatric surgery unit 1939-1971, and in 1957 published his influential “Textbook on Paediatric Surgery” with contributing authors Margrit Stockmann (Luzern), and Marcel Bettex, then consulting paediatric surgeon in Zurich. Grob’s “Textbook”, indiscriminately hailed by the Zurich University Children’s Hospital till this day, stressed the “special importance” for surgeons of Andrea Prader’s newly developed systematic classification of “genital variations” (“Prader Scales”). In its section on “surgical correction of the external genital” of children with 46,XX CAH (“[T]he removal of the enlarged clitoris [...] suggests itself. [...] Technique: [...] Usually we leave a very short clitoris stump”), Grob proclaimed the psychosocial justifications for cosmetic genital surgery on intersex children still prevalent today “The amputation of the clitoris, which may appear bothersome due to its size and erections, and may lead to embarrassment for these girls in the changing room or while swimming, is surely justified.” Grob became the founder and first president of the Swiss Society for Paediatric Surgery, and honorary member of the German, Austrian, British and U.S. societies. Grob’s recommendations in the “Textbook” (“surgical correction” in case of Prader Stages II–V, arguably devised at least with input by Prader himself), represented the global standard until the “Chicago DSD Consensus Conference” in 2005 (changing it to III–V).

Plastic Operations on the Genitalia

The **surgical correction** (see p. 474 *et seq.*) of the masculinized genitalia of girls with the congenital adrenogenital syndrome is desirable for several reasons: (1) in order to make the vagina a functional organ; (2) in order to prevent troublesome erections of the clitoris; (3) in order to prevent psychological conflicts, which are particularly liable to occur in girls with male characteristics.

Whenever possible surgery should be carried out **before the children reach four years of age**. In mild cases removal of the clitoris is all that is necessary. The clitoris should be **totally** removed and not just amputated, otherwise troublesome erections of the remaining stump may occur. As Hampson (1956) was able to show in a large series of women subjected to operation, removal of the clitoris does not interfere with the ability to achieve orgasm. If masculinization of the genitalia is more extreme further surgery may be required to open and enlarge the urogenital sinus.

**Source:**
Jürgen R. Bierich: “The Adrenogenital Syndrome”

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**1956–1993: “The Clitoris is not essential for normal Coitus.”**
“**No Evidence of Loss of Orgasm after Clitoris Amputation.”**

The number of “Intersex-Experts” and involved clinicians claiming that amputating “enlarged” clitorises was a rational and beneficent thing to do is legion – e.g. Joan Hampson (1956), John Money (1956, 1971), Max Grob (1957, see above), Jürgen Bierich (1963, 1971), Robert E. Gross (1966), Marcel Bettes (1957, see above).

Even in 1993, surgeon Milton Edgerton claimed, unchallenged by his peers: “Not one has complained of loss of sensation, even when the entire clitoris was removed.”

**Since then: “Surgery is better now ...”**

In 1993, Cheryl Chase founded the first Intersex Lobby Group ISNA by declaring: “Unfortunately the surgery is immensely destructive of sexual sensation and of the sense of bodily integrity.” Since then, the mutilators just changed their mantra to “Surgery is better now” – again without evidence, but despite survivors deploring decrease or total loss of sexual sensation, painful scars and frequent complications also with the “modern improved techniques”, and studies again and again corroborating their grievances.
