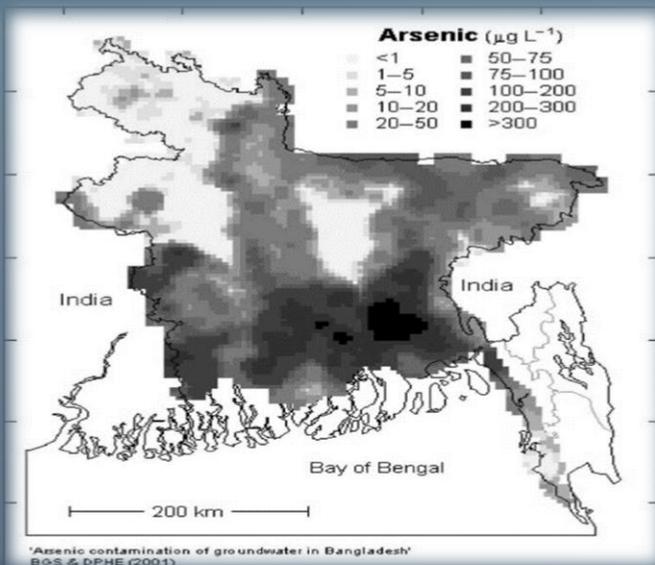


SEA LEVEL RISE AFFECTS GROUNWATER ARSENIC CONCENTRATIONS IN FOUR COASTAL DIVISIONS OF THE GANGES DELTA: IMPLICATIONS FOR WATER QUALITY SUSTAINABILITY

-MODELLED FROM THE YEAR 2000 UNTIL THE YEAR 2100-



Utrecht University



2050 sea level

2030 sea level

As Arsenic 74.922

30.974

33

51

50

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

60

61

62

63

64

65

66

67

68

69

70

71

72

73

74

75

76

77

78

79

80

81

82

83

84

85

86

87

88

89

90

91

92

93

94

95

96

97

98

99

100

101

102

103

104

105

106

107

108

109

110

111

112

113

114

115

116

117

118

119

120

121

122

123

124

125

126

127

128

129

130

131

132

133

134

135

136

137

138

139

140

141

142

143

144

145

146

147

148

149

150

151

152

153

154

155

156

157

158

159

160

161

162

163

164

165

166

167

168

169

170

171

172

173

174

175

176

177

178

179

180

181

182

183

184

185

186

187

188

189

190

191

192

193

194

195

196

197

198

199

200

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

217

218

219

220

221

222

223

224

225

226

227

228

229

230

231

232

233

234

235

236

237

238

239

240

241

242

243

244

245

246

247

248

249

250

251

252

253

254

255

256

257

258

259

260

261

262

263

264

265

266

267

268

269

270

271

272

273

274

275

276

277

278

279

280

281

282

283

284

285

286

287

288

289

290

291

292

293

294

295

296

297

298

299

300

301

302

303

304

305

306

307

308

309

310

311

312

313

314

315

316

317

318

319

320

321

322

323

324

325

326

327

328

329

330

331

332

333

334

335

336

337

338

339

340

341

342

343

344

345

346

347

348

349

350

351

352

353

354

355

356

357

358

359

360

361

362

363

364

365

366

367

368

369

370

371

372

373

374

375

376

377

378

379

380

381

382

383

384

385

386

387

388

389

390

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

408

409

410

411

412

413

414

415

416

417

418

419

420

421

422

423

424

425

426

427

428

429

430

431

432

433

434

435

436

437

438

439

440

441

442

443

444

445

446

447

448

449

450

451

452

453

454

455

456

457

458

459

460

461

462

463

464

465

466

467

468

469

470

471

472

473

474

475

476

477

478

479

480

481

482

483

484

485

486

487

488

489

490

491

492

493

494

495

496

497

498

499

500

501

502

503

504

505

506

507

508

509

510

511

512

513

514

515

516

517

518

519

520

521

522

523

524

525

526

527

528

529

530

531

532

533

534

535

536

537

538

539

540

541

542

543

544

545

546

547

548

549

550

551

552

553

554

555

556

557

558

559

560

561

562

563

564

565

566

567

568

569

570

571

572

573

574

575

576

577

578

579

580

581

582

583

584

585

586

587

588

589

590

591

592

593

594

595

596

597

598

599

600

601

602

603

604

605

606

607

608

609

610

611

612

613

614

615

616

617

618

619

620

621

622

623

624

625

626

627

628

629

630

631

632

633

634

635

636

637

638

639

640

641

642

643

644

645

646

647

648

649

650

651

652

653

654

655

656

657

658

659

660

661

662

663

664

665

666

667

668

669

670

671

672

673

674

675

676

677

678

679

680

681

682

683

684

685

686

687

688

689

690

691

692

693

694

695

696

697

698

699

700

701

702

703

704

705

706

707

708

709

710

711

712

713

714

715

716

717

718

719

720

721

722

723

724

725

726

727

728

729

730

731

732

733

734

735

736

737

738

739

740

741

742

743

744

745

746

747

748

749

750

751

752

753

754

755

756

757

758

759

760

761

762

763

764

765

766

767

768

769

770

771

772

773

774

775

776

777

778

779

780

781

782

783

784

785

786

787

788

789

790

791

792

793

794

795

796

797

798

799

800

801

802

803

804

805

806

807

808

809

810

811

812

813

814

815

816

817

818

819

820

821

822

823

824

825

826

827

828

829

830

831

832

833

834

835

836

837

838

839

840

841

842

843

844

845

846

847

848

849

850

851

852

853

854

855

856

857

858

859

860

861

862

863

864

865

866

867

868

869

870

871

872

873

874

875

876

877

878

879

880

881

882

883

884

885

886

887

888

889

890

891

892

893

894

895

896

897

898

899

900

901

902

903

904

905

906

907

908

909

910

911

912

913

914

915

916

917

918

919

920

921

922

923

924

925

926

927

928

929

930

931

932

933

934

935

936

937

938

939

940

941

942

943

944

945

946

947

948

949

950

951

952

953

954

955

956

957

958

959

960

961

962

963

964

965

966

967

968

969

970

971

972

973

974

975

976

977

978

979

980

981

982

983

984

985

986

987

988

989

990

991

992

993

994

995

996

997

998

999

1000

1001

1002

1003

1004

1005

1006

1007

1008

1009

1010

1011

1012

1013

1014

1015

1016

1017

1018

1019

1020

1021

1022

1023

1024

1025

1026

1027

1028

1029

1030

1031

1032

1033

1034

1035

1036

1037

1038

1039

1040

1041

1042

1043

1044

1045

1046

1047

1048

1049

1050

1051

1052

1053

1054

1055

1056

1057

1058

1059

1060

1061

1062

1063

1064

1065

1066

1067

1068

1069

1070

1071

1072

1073

1074

1075

1076

1077

1078

1079

1080

1081

1082

1083

1084

1085

1086

1087

1088

1089

1090

1091

1092

1093

1094

1095

1096

1097

1098

1099

1100

1101

1102

1103

1104

1105

1106

1107

1108

1109

1110

1111

1112

1113

1114

1115

1116

1117

1118

1119

1120

1121

1122

1123

1124

1125

1126

1127

1128

1129

1130

1131

1132

1133

1134

1135

1136

1137

1138

1139

1140

1141

1142

1143

1144

1145

1146

1147

1148

1149

1150

1151

1152

1153

1154

1155

1156

1157

1158

1159

1160

1161

1162

1163

1164

1165

1166

1167

1168

1169

1170

1171

1172

1173

1174

1175

1176

1177

1178

1179

1180

1181

1182

1183

1184

1185

1186

1187

1188

1189

1190

1191

1192

1193

1194

1195

1196

1197

1198

1199

1200

1201

1202

1203

1204

1205

1206

1207

1208

1209

1210

1211

1212

1213

1214

1215

1216

1217

1218

1219

1220

1221

1222

1223

1224

1225

1226

1227

1228

1229

1230

1231

1232

1233

1234

1235

1236

1237

1238

1239

1240

1241

1242

1243

1244

1245

1246

1247

1248

1249

1250

1251

1252

1253

1254

1255

1256

1257

1258

1259

1260

1261

1262

1263

1264

1265

1266

1267

1268

1269

1270

1271

1272

1273

1274

1275

1276

1277

1278

1279

1280

1281

1282

1283

1284

1285

1286

1287

1288

1289

1290

1291

1292

1293

1294

1295

1296

1297

1298

1299

1300

1301

1302

1303

1304

1305

1306

1307

1308

1309

1310

1311

1312

1313

1314

1315

1316

1317

1318

1319

1320

1321

1322

1323

1324

1325

1326

1327

1328

1329

1330

1331

1332

1333

1334

1335

1336

1337

1338

1339

1340

1341

1342

1343

1344

1345

1346

1347

1348

1349

1350

1351

1352

1353

1354

1355

1356

1357

1358

1359

1360

1361

1362

1363

1364

1365

1366

1367

1368

1369

1370

1371

1372

1373

1374

1375

1376

1377

1378

1379

1380

1381

1382

1383

1384

1385

1386

1387

1388

1389

1390

1391

1392

1393

1394

1395

1396

1397

1398

1399

1400

1401

1402

1403

1404

1405

1406

1407

1408

1409

1410

1411

1412

1413

1414

1415

1416

1417

1418

1419

1420

1421

1422

1423

1424

1425

1426

1427

1428

1429

1430

1431

1432

1433

1434

1435

1436

1437

1438

Table of contents

Summary	3
Introduction	3
Concepts and theories	5
Concepts	5
Theories	7
Literature review	10
Conceptual framework	11
Analytical framework.....	12
Methodology	13
Methodology first sub question	13
Methodology second sub question	13
Methodology third sub question and main research question	13
Description of study area.....	13
Data selection	14
Validation, replicability, replication and justification	14
Data analysis.....	15
Ethical issues	15
Results	16
Sub question 1	16
Sub question 2	18
Sub question 3	18
Main research question.....	19
Discussion	20
Answering the sub questions and main research question	20
Compare findings to other research – validation.....	21
Limitations and strengths	22
Implications, further research and recommendations	23
Conclusion	24
References	25
Appendix I: Additional calculations and explanation formulas	29
Calculation of the study area	29
Starting molarity bicarbonate	29
Starting molarity CO ₂	30
Relation pH and redox potential.....	30
Relation between arsenic and iron hydroxide (Fe ²⁺)	30

Relation between iron hydroxide (Fe^{2+}) and redox potential	30
Relationship between HCO_3^- and iron hydroxide.....	30
Relation pH and arsenic	30
Relation flux bicarbonate and CO_2	30
Appendix II: The model in Excel	31
Appendix III: Formulas and variables clarified.....	32
Appendix IIII: Other graphs for data analysis derived from SPSS	34
Arsenic concentrations derived from the pH.....	34
Arsenic concentrations derived from redox potential.....	34
Arsenic concentrations derived from iron hydroxide	35
Multiple regression analysis	36
Descriptive statistics.....	37

Summary

In Bangladesh, and especially in the groundwater of the four coastal divisions of the Ganges delta, the arsenic concentrations are above the limits set by the WHO and Bangladesh, which causes the population to experience severe health effects. Bangladesh is also susceptible to climate change, as it is heavily affected by the sea level rise and the accompanied salt water intrusion. This will change the conditions in the groundwater of the Ganges delta and subsequently affect the arsenic concentrations, but it is still unknown how, since no link has yet been found between these factors. The research question therefore is: *How will the effective sea level rise affect arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta from the year 2000 until the year 2100?* To research this question, Excel and SPSS are used and a literature review is conducted. Excel models the change of arsenic concentrations over time by using formulas found through the literature review. These formulas are analysed in SPSS. The results show that the effective sea level rise in Bangladesh amounts to 0.35 m between 2000 and 2100 and that arsenic concentrations will additionally change due to their relation with the molarity of bicarbonate and CO₂, pH, redox potential and ferrous iron. The molarity of bicarbonate and CO₂ will increase from 2000 to 2100 by 2.5 mol/L and by 0.03 mol/L. This subsequently increases the pH by 0.4, which directly influences the arsenic concentrations. The pH also has an indirect influence on arsenic through the redox potential and ferrous iron. The redox potential will decrease by 55 mV and ferrous iron will increase by 4.5 mg/L from 2000 to 2100. Together, the indirect and direct effect increase the arsenic concentration by 317 µg/L with a value of 1017 µg/L by 2100. Implications from this research are that the knowledge gap between sea level rise and the hereby changing arsenic concentrations is closed and that long-term predictions about arsenic concentrations can be done. This leads to the recommendation of implementing more long term measures to reduce the arsenic concentrations in groundwater.

Introduction

In the delta of Bangladesh, also known as the Ganges delta, high arsenic concentrations occur in the aquifers and groundwater (Nickson et al., 1998). This is especially the case in the four coastal divisions of the Ganges delta, which are Khulna, Barisal, Comilla and Chittagong (Appendix I; figure 1). These high arsenic concentrations are above the limits set by the World Health Organisation, which is 10 µg/L, and by Bangladesh, which is 50 µg/L (Smith, Lopipero, Bates & Steinmaus, 2002). These high arsenic concentrations have severe health effects on the inhabitants of Bangladesh. Already 33 to 77 million people are affected and have developed certain types of cancer, skin pigmentation and keratosis (Smith, Lingas & Rahman, 2000). This is not the only problem that occurs in the Ganges delta, as Bangladesh is considered to be the most vulnerable country to climate change (Paul & Rashid, 2017). The Ganges delta is heavily affected by sea level rise, and experiences frequent flooding (Karim & Mimura, 2008). This flooding causes salt water intrusion, which affects the groundwater conditions in the Ganges delta. These changed conditions subsequently affect the arsenic concentrations in this area (Snyder, Taillefert & Ruppel, 2004; Nordstrom, 2002). Therefore, the research question is: *How will the effective sea level rise affect arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta from the year 2000 until the year 2100?* To answer the research question, three sub questions have been formed. The first sub question is: *Which factors are influencing and causing the high arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta?* The second sub question is: *How much will the*

effective sea level rise be in the Ganges delta from the year 2000 until the year 2100? The third sub question is: How will the rising sea level change these factors and eventually thus the arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta?

It is currently known that the rising sea level increases the amount of salt water intrusion (Paul & Rashid, 2017) and it is known that arsenic concentrations are high (Anawar et al., 2003). However there is a gap between these two issues, since it is still unknown how the sea level rise and the hereby changing groundwater conditions will influence the arsenic concentrations in the Ganges delta. Additionally, it is unknown how the population of Bangladesh will be affected by the arsenic concentrations in the future. It is for both the society and the government of Bangladesh of great importance to know whether and how the arsenic concentrations will change due to the rising sea level, because this way they know whether they need to implement more or less measures and technical solutions in the future to reduce the amount of arsenic in the groundwater. This again will reduce the amount of people affected. This also indicates the connection of this research to sustainability; researching changes in arsenic concentrations due to the effective sea level rise is of great importance, because research needs to be done on how this implicates the water quality sustainability. Furthermore, sustainable solutions need to be developed to ensure access to safe drinking water for future generations.

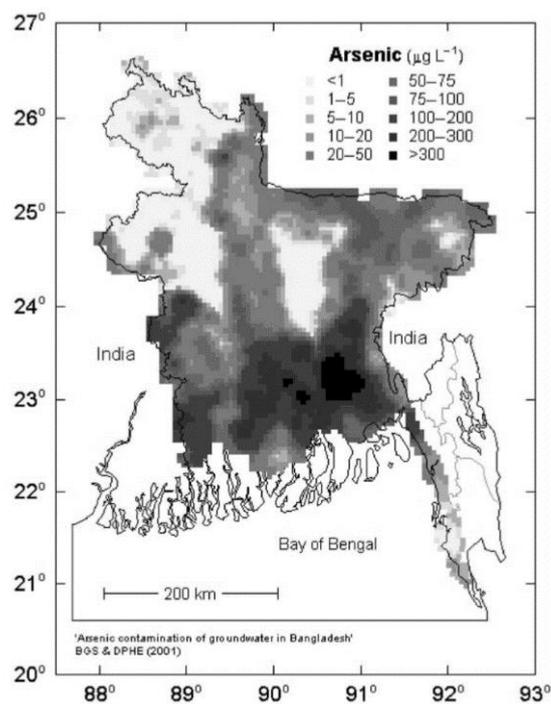


Fig. 1: Map showing the concentrations of arsenic in Bangladesh, which increase towards the coast (Ahmed et al., 2004).

Concepts and theories

Concepts

There are several key concepts related to this research, which are described and explained below.

Eustatic sea level rise

The eustatic sea level rise is determined by the volume of the global ocean and resembles the absolute sea level rise. It can increase due to melting of glaciers and ice caps and due to thermal expansion (Pethick & Orford, 2013; Paul & Rashid, 2017; Kay et al., 2015). Thermal expansion is the phenomenon whereby seawater increases in temperature, which decreases the density (Shamsuddoha & Chowdhury, 2007). This is currently happening, due to global warming (Paul & Rashid, 2017).

Subsidence

Subsidence is the phenomenon whereby the surface sinks (Brammer, 2014). It is caused by several processes, which are tectonics, human influences, sedimentation and sediment compaction. In Bangladesh, tectonics have a big influence on subsidence, but scientists also believe that compaction plays a large role in the subsidence of the Ganges delta (Brown & Nicholls, 2015; Paul & Rashid, 2017). Subsidence is approximately 2.0 mm/year, but according to other research can be as high as 20 mm/year (Brammer, 2014; Paul & Rashid, 2017).

Tectonic uplift

Tectonic uplift is a phenomenon whereby the surface rises. This is due to past ice ages, when water was stored in large glaciers, which made the surface subside. Currently, there is an interglacial period, in which the glaciers melt and pressure is released. This causes the uplift of the surface (Paul & Rashid, 2017). In Bangladesh, tectonic uplift lifts the surface by 3.6 mm/year (Paul & Rashid, 2017).

The effective sea level rise

The effective sea level rise resembles the relative sea level rise, which is the rate of the sea level change relative to the delta surface and is influenced by the level of the delta surface and the eustatic sea level rise. The level of the delta surface can change because of subsidence and tectonic uplift, which are described above (Pethick & Orford, 2013; Karim & Mimura, 2008; Shamsuddoha & Chowdhury, 2007; Paul & Rashid, 2017).

Salt water intrusion

Because of the sea level rise, more and further-inland intrusion of seawater will occur in the future (Ministry of foreign affairs, 2018; Pethick & Orford, 2013). This salinization is secondary salinization, since the increase in salt concentrations is not due to the change between glacial and interglacial cycles (primary salinization), but due to global climate change (Paul & Rashid, 2017). Seawater has a different composition than regular saline water. Seawater contains mostly Sodium (Na^+), Potassium (K^+), Calcium (Ca^{2+}), Magnesium (Mg^{2+}), Chloride (Cl^-), Carbonate (CO_3^-), Bicarbonate (HCO_3^-) and sulphate (SO_4^-) (Millero, Feistel, Wright & McDougall, 2008; Paul & Rashid, 2017).

Arsenic

Arsenic (As) is a chemical element (number 33 in the periodic table) that can be taken up into the minerals of other elements, such as pyrite (FeS). Once arsenic is taken up into the mineral of pyrite, pyrite changes into arseno-pyrite (AsFeS). Arsenic can also be adsorbed onto other elements, such as iron and can be in competition for adsorption places with other elements, such as phosphorus (Van der Perk, 2013). Arsenic occurs naturally in the groundwater of Bangladesh and the concentrations of arsenic here are abnormally high, since the concentrations are far above the limits set by both the World Health Organisation, 10 µg/L, and Bangladesh, 50 µg/L (Smith et al., 2002; Nickson et al., 1998).

pH

The pH is the hydrogen ion concentration (H^+) in a solution. This can be calculated using the following formula: $pH = -\log[H^+]$. When the pH is below 7, the solution has a high concentration of H^+ ions and the solution is acidic. At pH is 7, the solution is neutral and when the pH is above 7, the solution is alkaline. This means that the solution has few H^+ ions and high concentrations of OH^- ions (Hendriks, 2010; Hem, 1972).

Redox potential

Redox potential (E_h) is a concept that is involved in redox processes. Redox processes are chemical reactions, in which an electron from one ion is transferred to another ion. The redox potential is the tendency of an ion to lose or acquire an electron in a redox process, which makes the ion reduced or oxidised (Hem, 1972; Van der Vecht & Gijben, 2016). The redox reaction rates and directions are determined by the lack of the availability of oxygen in the environment. Oxygen has the highest redox potential and is therefore first in line to be reduced to H_2O (see fig. 2). Organic matter has the lowest redox potential and is the ultimate reductant. Hence, organic matter consumes oxygen, which leads to bacterial reduction and the breaking down of organic matter (Groundwater Geochemistry, n.d.). However, when oxygen is not available, another substance is needed to break down organic matter through microbial degradation. Then, the order of redox potential is followed (see figure 2). When oxygen is depleted, nitrate is used as a reducer instead of oxygen and so forth (Van der Perk, 2013; Groundwater Geochemistry, n.d.).

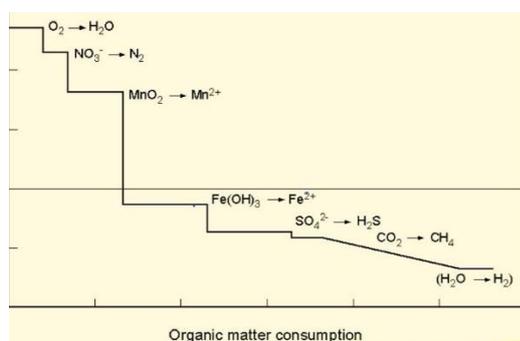


Fig. 2: Redox reactions; the lower on the ladder, the lower the potential (Van der Perk, 2013).

Theories

The theories described below are connecting the mechanisms of arsenic mobilisation to the concepts explained above.

Reduction hypothesis

The most supported hypothesis about arsenic mobilisation is the reduction hypothesis (Ahmed et al., 2004). The reduction hypothesis is connected to the concept redox potential, which is explained above. When oxygen, nitrate and manganese are all depleted, ferric iron ($\text{Fe}(\text{OH})_3$) can be used as a reducer to break down organic matter (Van der Perk, 2013). In an oxic environment, arsenic species can adsorb onto iron hydroxide by replacing a hydroxide ion (H^+) on the surface of iron hydroxide (Van der Perk, 2013; Anawar et al., 2003; Jain, Raven & Loeppert, 1999). In an anoxic environment, ferric iron is reduced and is transferred into ferrous iron (Fe^{2+}) as is displayed in figure 2. Ferrous iron is a form of iron that is much more soluble than ferric iron (Anawar et al., 2003). Because of this increased solubility, arsenic cannot readily be adsorbed onto the surface and is released (Brammer & Ravenscroft, 2009). This is what is meant by the reduction hypothesis: the reductive dissolution of arsenic-rich iron (Nickson, McArthur, Ravenscroft, Burgess & Ahmed, 2000).

pH in connection with arsenic

The pH is also connected to arsenic, which can readily be explained when looking at the mechanism of deprotonation. When the water is more alkaline, H^+ ions are removed (deprotonation) from the surface of iron hydroxide due to competition with the other bases in solution. As a consequence, the negative charge on the surface of the iron hydroxides increases with increasing pH and thus the capacity to adsorb anions, such as arsenate/arsenite decreases. So the higher the pH, the more arsenic is mobilised and the higher the concentrations in the groundwater (Van der Perk, 2013; Anawar et al., 2003).

Salt water intrusion and pH

Salt water intrusion into groundwater leads to the mixing of seawater and fresh water. This causes a change in the pH, which again has an effect on arsenic, as is described above (Anawar et al., 2003). The pH is influenced by the concentration of H^+ ions in the water. The amount of CO_2 and bicarbonate (HCO_3^-) in the water influences the amount of H^+ ions. The increase of CO_2 in the seawater due to climate change will not be taken into account, since this goes beyond the scope of this study. The relation between CO_2 and H^+ is displayed in this equilibrium formula:



H_3O^+ is similar to H^+ , as it is a H^+ ion that has reacted with water. Because the formula is an equilibrium formula, there is an acid dissociation constant, K_a . This is the equilibrium constant for an acid-base reaction (Van der Vecht & Gijben, 2016). For seawater, all the concentrations are known, so the K_a value can be calculated using the equilibrium formula and the pH-formula (Van der Vecht & Gijben, 2016; Hendriks, 2010):

$$K_a = \frac{[\text{H}_3\text{O}^+][\text{HCO}_3^-]}{[\text{CO}_2]} \quad \text{pH} = -\log[\text{H}^+] \rightarrow \text{H}^+ = 10^{-\text{pH}}$$

It is known that the pH of seawater is 8.1, which will be used to calculate the molarity of H^+ in mol/L: $H^+=10^{-8.1} = 7.94 \times 10^{-9}$ mol/L (Millero et al., 2008). The value of $[HCO_3^-]$ is known in mol/kg, which is 0.0017803. It can be recalculated to mol/L by using the density of seawater. The density of seawater is 1.025 kg/L, so $0.0017803 \times 1.025 = 1.83 \times 10^{-3}$ mol/L (Millero et al., 2008; Hendriks, 2010). $[CO_2]$ value is also known in mol/kg, which is 0.0000100 mol/kg and is transferred to mol/L by again using the density of seawater: $0.0000100 \times 1.025 = 1.025 \times 10^{-5}$ mol/L (Millero et al., 2008).

Inserting these numbers into the formula gives

$$K_a = \frac{[7.94 \times 10^{-9}][1.83 \times 10^{-3}]}{[1.025 \times 10^{-5}]} = 1.414 \times 10^{-6} \text{ (Millero et al., 2008; Van der Vecht \& Gijben, 2016).}$$

Because this K_a -value stays the same for the reaction in groundwater, and the molarities in groundwater will be calculated for CO_2 and HCO_3^- , the pH can be calculated for groundwater.

Relation of pH and redox potential

A change in redox potential (E_h) is related to the pH of the surrounding water. As was described above, the pH will change when seawater and groundwater mix. Due to this mixing, the pH of the groundwater will increase. The higher the pH, the lower the redox potential (Xyla et al., 1992). This is indicated by the Nernst equation:

$$E_h = E_0 + \frac{0,05916}{n} \log \left(\frac{\{A\}\{B\}}{\{C\}\{D\}} \right) - \frac{0,05916h}{n} pH \text{ (Bundschuh \& Sracek, 2011).}$$

When the pH increases by 1.5, the redox potential will decrease by 200 mV (Sondergaard, 2009). The lower the redox potential, the more an electron wants to be released, which causes the oxidation of the new species (Oregon State University, n.d.; Sondergaard, 2009). This means that there will be more ferric iron transferred to ferrous iron. Ferrous iron is more soluble and has less adsorption sites, which means that more arsenic desorbs (Sondergaard, 2009). An increase in pH and hereby decreased redox potential will thus both lead to higher arsenic concentrations.

The connection between sea level rise and groundwater

The rate of sea level rise has an influence on the amount of seawater that ends up in the groundwater. This again has an influence on the molarities of HCO_3^- and CO_2 in the groundwater and subsequently on the pH (Bozlee & Janebo, 2008). The influence of the rate of sea level rise on the amount of seawater in an aquifer/groundwater can be determined using the Ghijben-Herzberg relation. This is the relation between freshwater above the mean sea level and freshwater below sea level, which is similar to the amount of freshwater in an aquifer. The relation is this:

$$z = \frac{\rho_f}{\rho_s - \rho_f} h.$$

ρ_f =the density of freshwater, ρ_s =the density of seawater, z =the depth of the freshwater above sea level and h =the depth of the freshwater below sea level (Hendriks, 2010; Werner & Simmons, 2009). Currently, z is 6 meters, since the coastal areas of the Ganges delta are still 6 meters above sea level, according to figure 3 and Tareq, Safiullah, Anawar, Rahman & Ishizuka

(2003). This means that there is still a recharge area of 6 m. However, this will get smaller due to sea level rise, which also results in a decrease of h . The decrease in h will reveal the area of groundwater that is affected by the sea level rise.

When this is determined, the new molarities of bicarbonate and CO_2 in the groundwater due to salt water intrusion can be calculated using Fick's law of diffusion. Diffusion is the process in which substances move from a high concentration to a low concentration, which is due to the concentration gradient (Van der Perk, 2013). Seawater has a different concentration of substances than groundwater, so when seawater enters the groundwater, there are differences in the concentrations of bicarbonate and CO_2 . Because of these differences in concentration, diffusion will take place. In this research, molarity will be used instead of concentration. Fick's law of diffusion is displayed by this formula:

$$J = -D \frac{c_2 - c_1}{x_2 - x_1}$$

J =the flux of the substance in $\text{mol/m}^2/\text{s}$, D =the diffusion coefficient, which is $1.6 \cdot 10^{-9} \text{ m}^2/\text{s}$ for bicarbonate and $2.1 \cdot 10^{-9}$ for CO_2 , c_2/c_1 =the concentrations of bicarbonate/ CO_2 in seawater and fresh water or in this research the molarity of both bicarbonate and CO_2 and x_2/x_1 =the distance of diffusion, which in this case is the decrease in h (Van der Perk, 2013; Solutions to the diffusion equation, 2006; Zeebe, 2011).

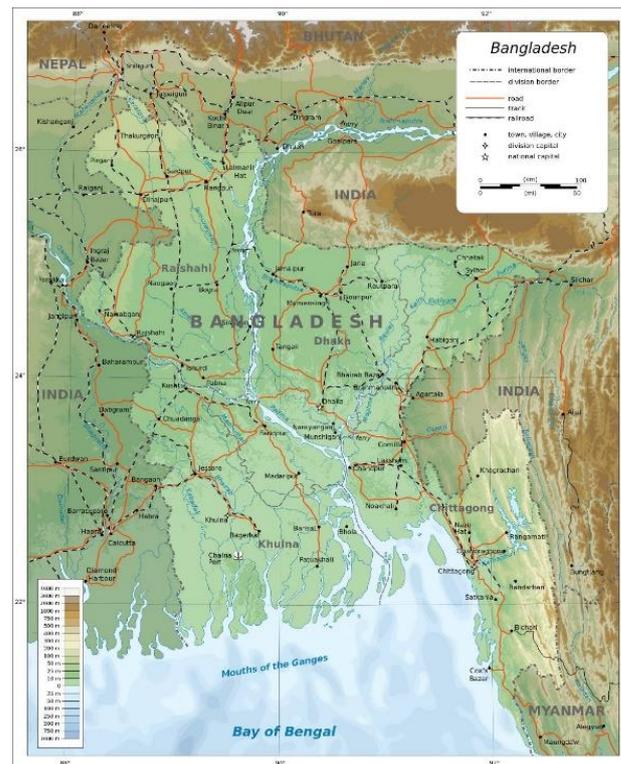


Fig. 3: Surface level relative to the sea level (Large detailed physical map of Bangladesh, n.d).

Literature review

In this research, literature is used which can be divided into three themes. The first theme is literature related to the mobilisation processes of arsenic (sub question 1). This literature was used to determine the most important variables related to the potential change of arsenic concentrations, such as redox potential and pH.

The second theme is related to the sea level rise (sub question 2), such as the tectonic uplift, subsidence and eustatic sea level rise. These variables were all found through the literature related to this theme and were the first input variables of the model.

The third and last theme is related to the connections of the variables found for the first theme described above. This theme is thus related to the redox potential, pH and other variables that were found to have an influence on the arsenic concentrations. This theme specifically looks at the formulas and connections between these variables to ensure that there is a connection and to ensure that the model works (sub question 3).

The division of literature into themes (see table 1) displays the contribution of these sources to the existing body of literature. There is a theme on mobilisation processes of arsenic and a theme related to the sea level rise. However, a third theme is needed to connect these themes, because there is a knowledge gap.

Table 1: This table displays which literature contributes to which theme.

THEME	LITERATURE CONTRIBUTING TO THEME
1. MOBILISATION PROCESSES OF ARSENIC	<ul style="list-style-type: none"> • Aggarwal et al., 2000 • Ahmed et al., 2004 • Anawar, 2003 • Anawar, 2002 • Brammer & Ravenscroft, 2009 • Jain et al., 1999 • Nickson et al., 1998 • Nickson et al., 2000 • Nordstrom, 2002 • Van der Perk, 2013
2. SEA LEVEL RISE	<ul style="list-style-type: none"> • Brammer, 2014 • Brown & Nicholls, 2015 • Karim & Mimura, 2008 • Kay et al., 2015 • Paul & Rashid, 2017 • Pethick & Orford, 2013 • Shamsuddoha & Chowdhury, 2007
3. CONNECTIONS BETWEEN VARIABLES	<ul style="list-style-type: none"> • Bhattacharya, Jacks, Ahmed, Routh & Khan, 2002 • Bozlee & Janebo, 2008 • Bundschu & Sracek, 2011 • Groundwater geochemistry, n.d. • Hem, 1972 • Hendriks, 2010 • Kvaternyuk et al., 2016 • Millero et al., 2008 • Nickson et al., 1998

	<ul style="list-style-type: none"> • Oregon State University, n.d. • Paul & Rashid, 2017 • Solutions to the diffusion equation, 2006 • Sondergaard, 2009 • Tareq et al., 2003 • Van der Perk, 2013 • Van der Vecht & Gijben, 2016 • Werner & Simmons, 2009 • Xyla et al., 1992 • Zeebe, 2011
<p>4. SOURCES USED FOR OTHER PURPOSES, SUCH AS INTRODUCTION, INFORMATION ON INPUT OF DATA AND INFORMATION ON EXCEL</p>	<ul style="list-style-type: none"> • Ahmed, 2001 • Baier & Neuwirth, 2003 • Bangladesh on the globe, 2014 • Billo, 2004 • Field 2018 • Large detailed map of Bangladesh, n.d. • Mahanta et al., 2015 • Ministry of foreign affairs, 2018 • Smith et al., 2000 • Smith et al., 2002 • Snyder et al., 20024 • Yang et al., 2014

Conceptual framework

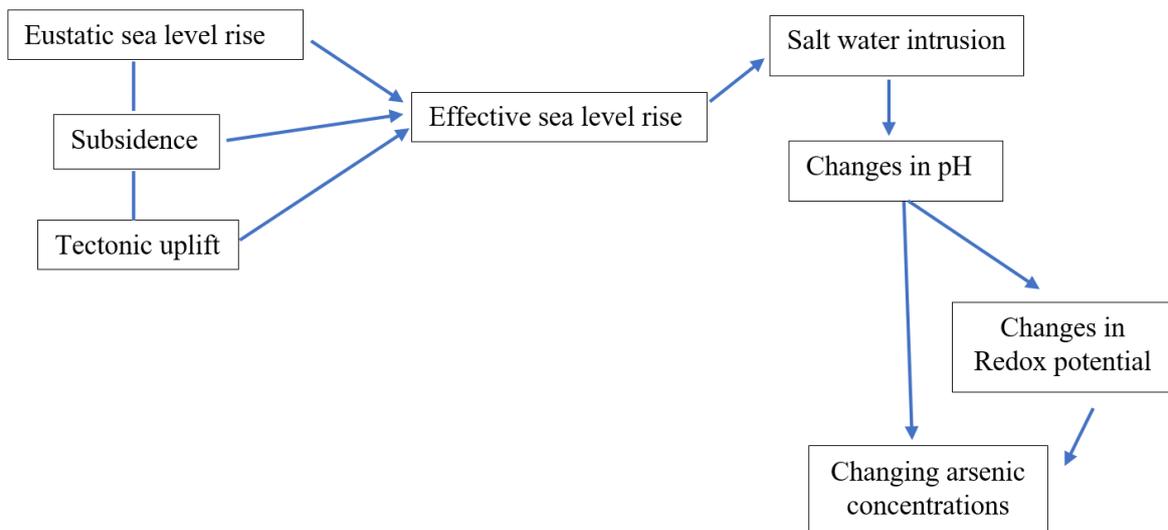


Fig. 4: The conceptual model gives an overview of the relations between the concepts and serves as the first step to visualising the model.

The conceptual framework displays the relations between the concepts. Eustatic sea level rise, subsidence and tectonic uplift all affect the effective sea level rise. The effective sea level rise has an effect on salt water intrusion, which is described under theories in ‘The connection between sea level rise and groundwater/aquifers’. Salt water intrusion has an effect on the pH, which has an effect on the redox potential. This is explained in theories under ‘Salt water intrusion and pH’ and ‘Relation of pH and redox potential’. The pH and redox potential both affect arsenic concentrations in the Ganges delta, which is explained in theories under ‘pH in connection with arsenic’ and ‘The reduction hypothesis’.

Analytical framework

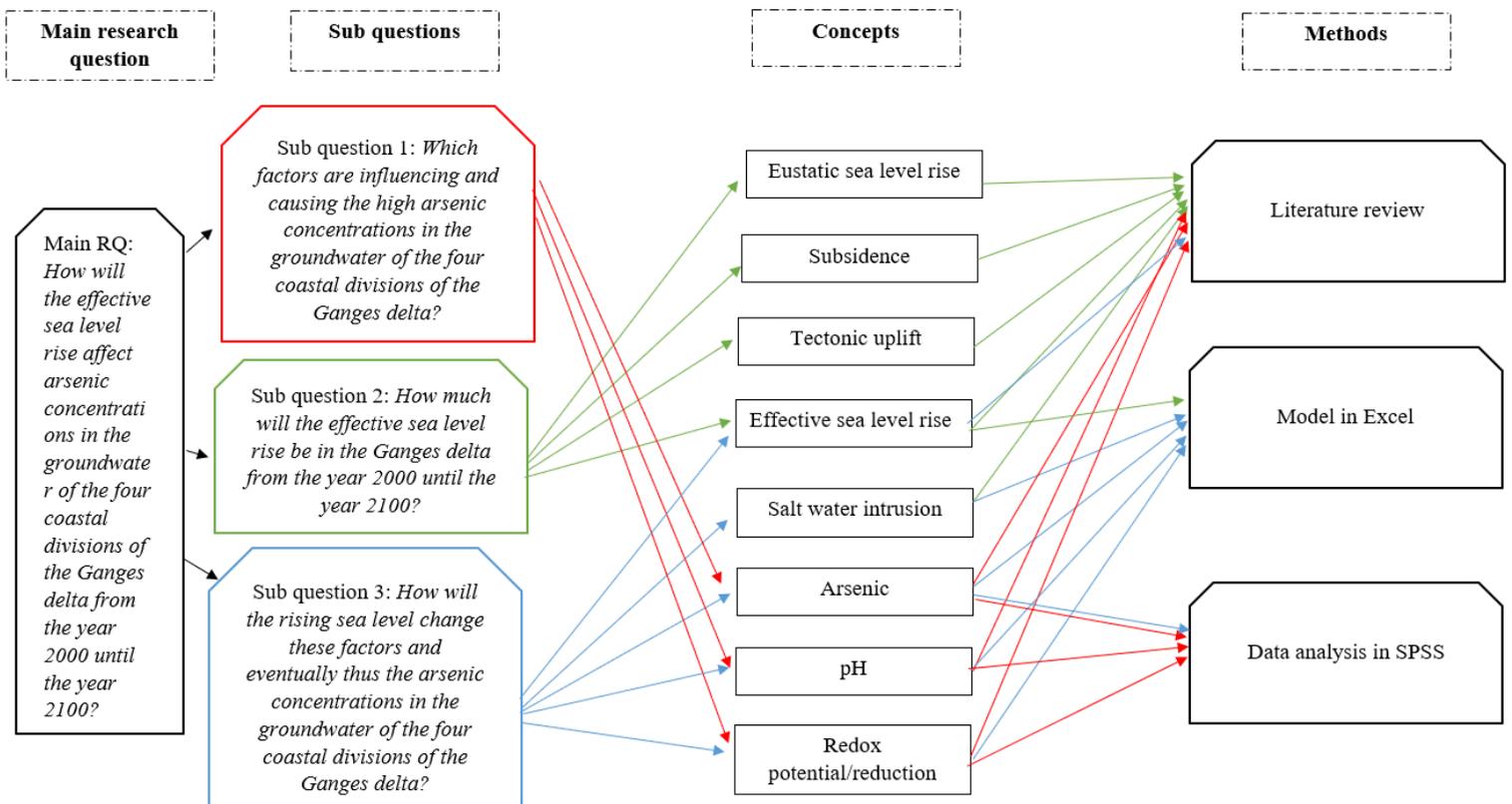


Fig. 5: The analytical model connects the sub questions, through all the concepts that were discussed, to the methods that are used in this research.

In figure 5, the analytical framework is displayed, which shows how the sub questions and related key concepts will be measured. It is divided into four parts: the main research question, the sub questions, the key concepts and the methods. In the method section, the analytical framework is explained in more detail.

Methodology

Methodology first sub question

In this research, there is one main research question and three sub questions which all require their own methodology. The first sub question relates to the factors that are influencing and causing the high arsenic concentrations (the red box in figure 5). For this sub question, a literature review must be conducted to determine which factors influence the mobilization processes of arsenic. In the described theories and concepts, two important factors are already identified to have the largest influence on arsenic concentrations, which are the pH and redox potential. Additionally, the relations between these factors need to be analysed in SPSS through regression analysis and a multiple regression analysis will be done to discover which of these factors influence the arsenic concentrations the most.

Methodology second sub question

The second sub question (the green box in figure 5) is related to the sea level concepts. For these concepts, such as subsidence, eustatic sea level rise and tectonic uplift, data are found through the literature review. Data collection is crucial for this sub question, since it is assumed that the pH has an empirical relation to the effective sea level rise. The effective sea level rise is modelled in Excel, which is feasible, since no yearly numbers are needed for the related variables, solely numbers that indicate the increase per year.

Methodology third sub question and main research question

To answer the third sub question (the blue box in figure 5), first a literature review needs to be conducted, which indicates how the sea level rise will change the factors that influence the arsenic concentrations and is already shortly described in ‘theories’. After data is found in the literature review for the changed conditions, the second part of sub question 3 and the main research question will be answered by building a model in Excel in which the data found in the literature review will be used. Excel will provide graphs, that visualise the changes of the factors that influence arsenic concentrations due to sea level rise. SPSS will visualise changes in arsenic concentrations in groundwater over time through descriptive statistics for the main research question.

Description of study area

The wider study area is Bangladesh (figure 6), which is a country in Southern Asia with a coastal area that covers 32% of the country’s area. About 46 million people live in this coastal area (Paul & Rashid, 2017). Bangladesh has a large delta, which forms due to deposition of sediments by a river that mouths into the ocean (Ministry of foreign affairs, 2018). The delta in Bangladesh is the largest delta in the world and is called the Ganges-Brahmaputra delta. It is adjacent to the Bay of Bengal (Shamsuddoha & Chowdhury, 2007) and consists of layers of silt, sand and clay. The sediments deposited by the Ganges rivers consist mostly of heavy clay (Aggarwal et al., 2000). The specific study area comprises of the four coastal divisions of the Ganges delta: Khulna, Barisal, Comilla and Chittagong (figure 7). These four divisions are adjacent to the Bay of Bengal and are all under the distance of the calculated 80 km (see appendix I).



Fig. 6: Location of Bangladesh on a world map (Bangladesh on the globe, 2014).

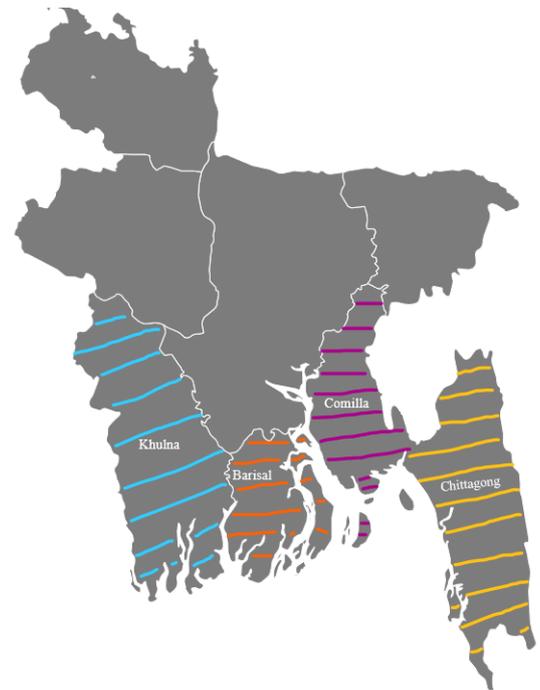


Fig. 7: Map showing the four coastal divisions of the Ganges delta: blue=Khulna; orange=Barisal; purple=Comilla and yellow=Chittagong.

Data selection

To clarify, three types of methods will be used in this bachelor thesis: a literature review, modelling in Excel and analysis through SPSS. For building the model in Excel, a chain of formulas is used which connects all the variables. These formulas are found through the literature review and will be analysed using SPSS to check their regression and significance. The type of data that is collected will be quantitative and no databases will be used. The data will be selected based on the published year of the article, preferably around the year 2000, since the timeline of this research extends from the year 2000 to 2100. The year 2100 is chosen as the last year, because this year is mostly used for predictions in climate. The data will additionally be selected based on location, as is described above under ‘description of study area’.

Validation, replicability, replication and justification

The literature review, Excel and SPSS are applicable in this research study and can be justified. The literature review is needed to form theories and find data and formulas that can be used to form the model and also to validate the model. Excel and SPSS can be justified by the short time frame for this research, combined with the acquaintance of the author with the software and accessibility of the software. Excel is easy to use and can be used to build a model that uses a chain of formulas for every separate year, but still provide the overall picture clearly, which the described problem in this research needs. Furthermore, Excel can organise large amounts of data into simple graphs and tables that are easy to read. Cells can also be connected, which means that it can adapt all of its data at once when new findings are found (Baier & Neuwirth, 2003).

Excel also has some shortcomings. It cannot check for human error when data is entered, which can have a strong negative influence on the results. Additionally, Excel can be time consuming when manually entering the data. In this research, the data is kept to a number that is feasible, so that mistakes are prevented (Billo, 2004).

There is a bulk of literature available about arsenic and sea level rise in which the variables involved in this research problem can be found. Yet, there are different ways to connect these variables in a model and different variables can be chosen from this literature. If another researcher would model this research, there is a possibility that different results would be found, which implicates the reliability slightly. The model will be validated in the discussion by comparing the found arsenic concentrations to the numbers and theories found in the literature. The model can be replicated using the provided formulas and data of this research. All the data will be provided in as much detail as possible, so that replication can be achieved (appendix III).

Data analysis

After the initial research and both before and after modelling in Excel, data analysis is conducted by using SPSS. This data analysis will focus on whether the formulas found in literature will have a high Pearson or Spearman correlation and whether these will produce significant results. If the formula seems correct, but another has a higher coefficient of determination, this formula will be used instead. To deal with uncertainty, a GLM ‘General Linear Model’ is performed in SPSS, which displays the Mean Squared Error and gives an indication of the uncertainty. The closer to zero, the lower the uncertainty. After the formulas are determined, a multiple regression analysis is performed to check which variable has the largest impact on changing the arsenic concentrations. This multiple regression analysis will be used to answer sub question 1 as is described above. The main research question also requires data analysis in the form of descriptive statistics. The change in arsenic concentrations will be analysed for every ten years, which displays in what period of time the change was largest.

Ethical issues

In this research, no field work or interviews are done, so only plagiarism and fraud are ethical issues that need to be prevented. Since the research problem is something that is never modelled before, fraud will be prevented. Plagiarism is avoided by citing the information used correctly, so that credit is given to the rightful authors. Shortly said, information that is written by other authors will be used but not copied, and credits will be given to these authors.

Results

The model is made in Excel through a chain of calculations (figure 15, appendix II). To do these calculations, formulas are needed, so the found formulas are analysed below under sub question 1 to confirm that there is a significant correlation. The other sub questions and main research question display graphs from after the model was built. Data analysis for these questions was only applicable to the main research question. The calculations done previous to the modelling part are displayed in appendix I. The results are structured by following the sub questions and main research question, since this corresponds to the analytical framework (figure 5).

Sub question 1

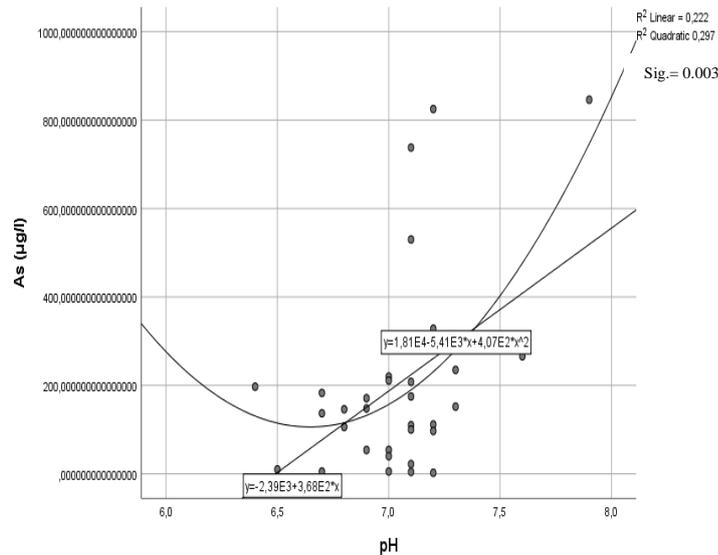


Fig. 8: Scatterplot of relation pH and arsenic concentrations with the coefficients of determination and significance in the right top corner.

The pH influences the arsenic concentrations, as can be seen in figure 8. In this figure, the relation between pH and arsenic is analysed. For the relation between pH and arsenic, a linear trend is chosen, since the difference in the coefficient of determination (R^2) is fairly small. Another reason to choose the linear trend is that the parabolic curve is very sensitive to specific scores and outliers and because the linear trend displays the main trend better (Field, 2018). The trend has a medium linear association (Field, 2018) and the correlation is significant according to the test of Pearson's R (table 6, appendix III). The ANOVA using GLM test indicates that the mean squared error, also known as the uncertainty, is still extremely large. However, since this is the formula with the highest R^2 and lowest uncertainty of the two formulas found, this formula is still used (table 7, appendix III). The linear trend displays that when the pH increases by 0.1, the arsenic concentrations increase by 36 $\mu\text{g/L}$. This is however an increase due to the direct relation of the pH and arsenic, and does not include the indirect effect due to the relation of pH, redox potential and ferrous iron described further below.

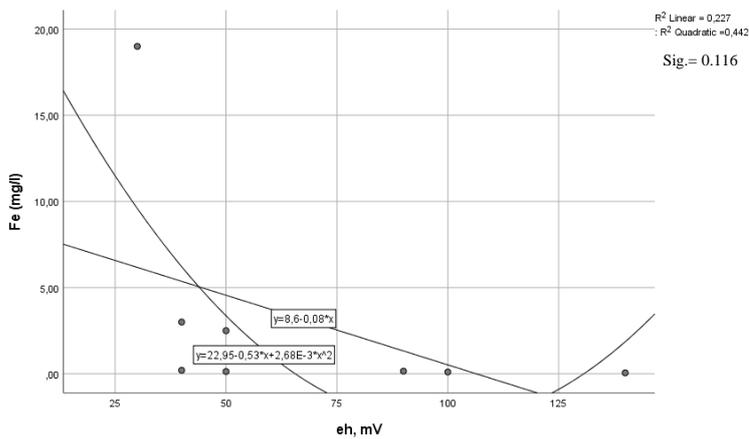


Fig. 9: Scatterplot of relation redox potential and ferrous iron with the coefficients of determination and significance in the right top corner.

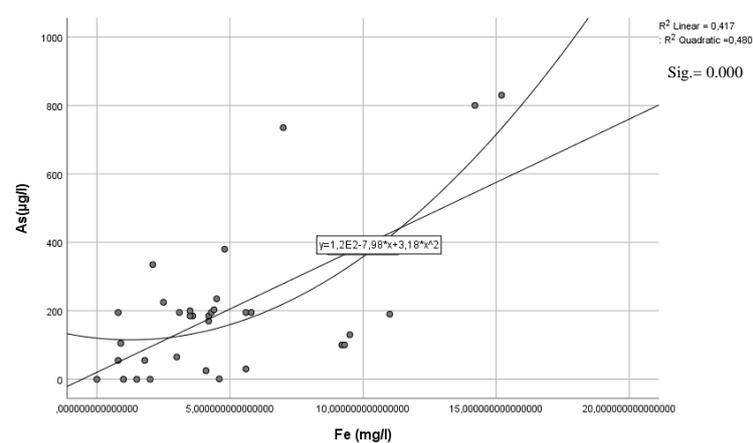


Fig. 10: Scatterplot of relation between ferrous iron and arsenic concentrations with the coefficients of determination and significance in the right top corner.

The pH, redox potential and ferrous iron are connected and affect the arsenic concentrations through a series of relations with each other. The formulas belonging to these relations are again analysed in figure 9 and 10. The relation between pH and redox potential is already perfectly linear, since there are only 4 data points, which is why data analysis is not conducted for this relation and the found formula is used (formula 12, appendix III).

The linear relation between the redox potential and the ferrous iron concentration has a coefficient of determination of 0.227 and the quadratic relation has a coefficient of determination of 0.442 (figure 9). Still the linear formula will be chosen to use in the model, since the quadratic trend has negative values between 125 and 75 mV, which is unrealistic. The linear line does explain the main trend and still has a medium linear association (Field, 2018). The Pearson's R test is again used for analysis, which shows a result that is not significant ($\text{sig.} = 0.116$). However, when bootstrapping is applied, the results are significant, since zero is not included in the confidence interval (table 8, appendix III). The formula will thus be used, especially because the ANOVA using GLM test shows that the mean squared error or the uncertainty is only 3.364, which is very small (table 9, appendix III).

Figure 10 describes the relationship between ferrous iron and arsenic. Again both quadratic and linear trends are analysed and the difference between the coefficients of determination is small. The linear trend is again chosen for the same reason as was described earlier. The coefficient of determination is 0.417, which indicates a strong linear association (Field, 2018). The Pearson's R test reveals a significance of 0.000, which means that the relation is significant. The uncertainty for this relation is 5909 (table 10 and 11, appendix III). The linear relation is positive and an increase of 1 mg/L will cause the arsenic concentrations to increase by about 37 µg/L (figure 10).

To test which of the three variables, the pH, redox potential and ferrous iron, has the largest influence on the arsenic concentrations, a multiple regression analysis is done. The multiple regression analysis shows that the pH has the highest standardised coefficient, so it has the largest influence on the arsenic concentrations. The redox potential has the second largest influence, since its standardised coefficient is the second largest. The ferrous iron concentration has the smallest influence on the arsenic concentrations, since its standardised coefficient is the smallest (table 12, appendix III).

Sub question 2

The model displays the change in effective sea level rise over the period 2000-2100. The relationship is linear and Excel displays that the sea level rise changes every year by 0.0035 m and by 0.35 m between 2000 and 2100.

Sub question 3

Change of ferrous iron and pH due to the effective sea level rise

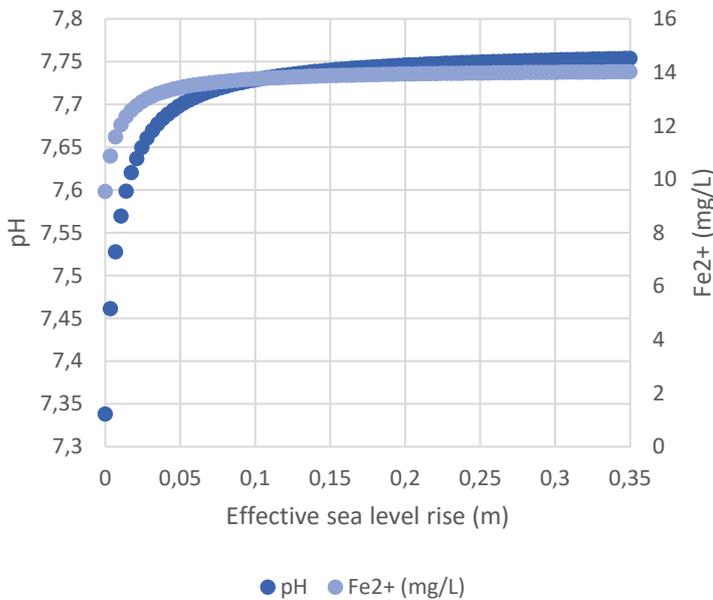


Fig. 11: The figure shows how the change in effective sea level rise changes the factors of pH and ferrous iron.

Change of molarity of H3O+ and redox potential due to the effective sea level rise

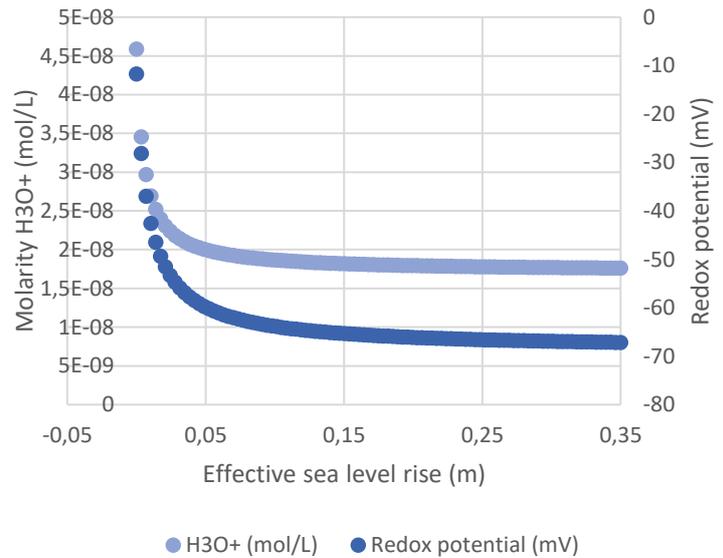


Fig. 12: The figure shows how the change in effective sea level rise changes the redox potential and the H₃O⁺ molarity.

Change of the molarity of bicarbonate and CO2 due to the effective sea level rise

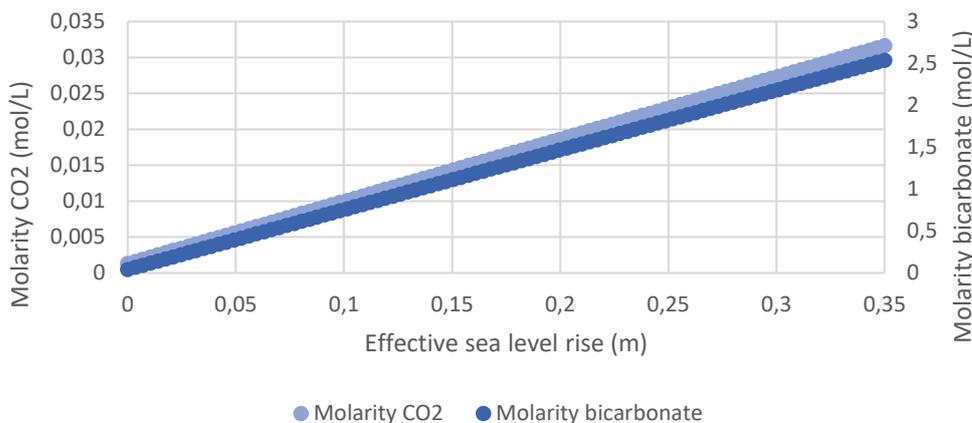


Fig. 13: The figure shows how the change in effective sea level rise changes the molarity of both CO₂ and HCO₃⁻.

The rising sea level will change the other variables through a chain of events. The higher the sea level rise, the higher the molarity of CO₂ and bicarbonate, the pH and the concentration of ferrous iron as can be seen in figure 11 and 13. An effective sea level rise of 0.35 m will increase

the molarity of bicarbonate by 2.5 mol/L, the molarity of CO₂ by 0.03 mol/L, the pH by about 0.4 and the concentration of ferrous iron by 4.5 mg/L. However, the higher the effective sea level rise, the lower the redox potential and the lower the molarity of H₃O⁺, as can be seen in figure 12. An effective sea level rise of 0.35 m will decrease the molarity of H₃O⁺ by about 2.8×10^{-8} mol/L and the redox potential by 55 mV. The relations of the CO₂ and bicarbonate molarities are linear to the sea level rise. The relations between the effective sea level rise and the pH, redox potential, the concentration of ferrous iron and the molarity of H₃O⁺ are however logarithmic.

Main research question

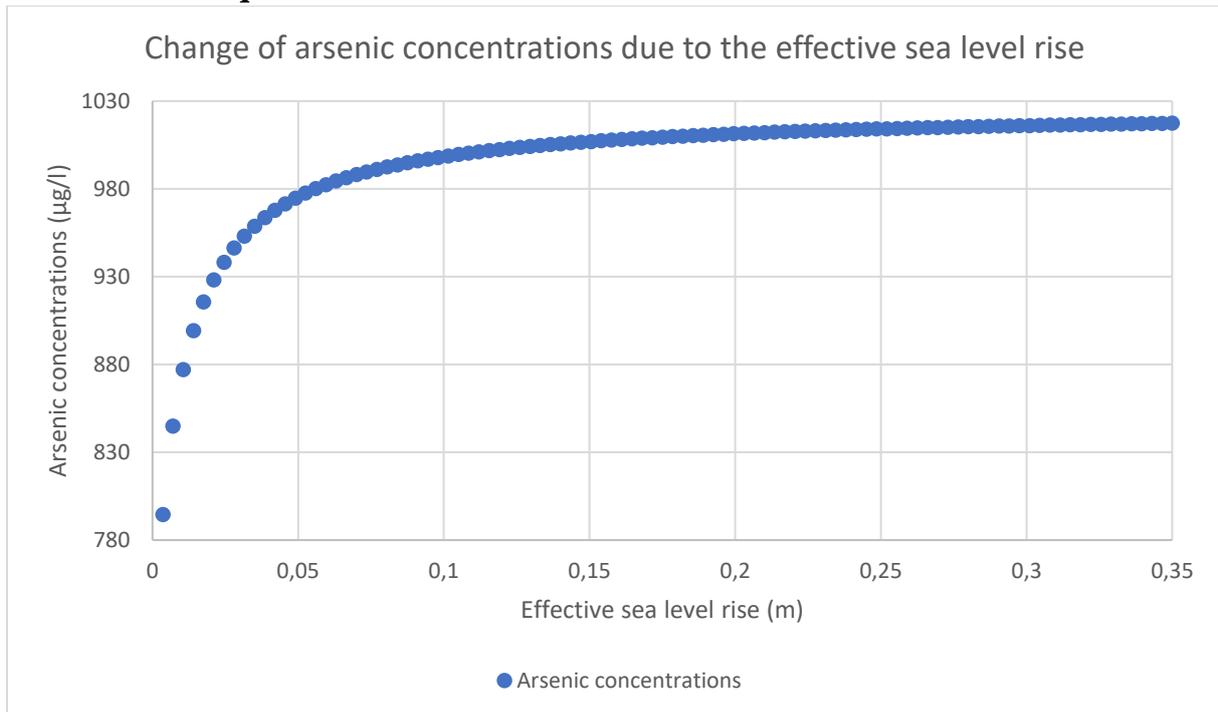


Fig. 14: The graphs shows how the effective sea level rise changes the arsenic concentrations.

Figure 14 shows that the effective sea level rise will cause higher arsenic concentrations through a logarithmic relationship. The graph displays that when the effective sea level rises by 0.35 m, the arsenic concentrations increase in total by 317 µg/L. In 2100, the arsenic concentrations amount to 1017 µg/L. The increase of arsenic concentrations slows down when the effective sea level rise increases and is different for different periods between 2000 and 2100, which is analysed using descriptive statistics (tables 13 through 22, appendix III). In these tables it is visible that the difference between the minimum and maximum value becomes smaller every ten years and that also every ten years the standard deviation, variance and standard error become smaller.

Discussion

Answering the sub questions and main research question

Research has been done on the arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta and how these concentrations would change due to the effective sea level rise. The research questions will be answered in this section.

The first sub question was: *Which factors are influencing and causing the high arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta?* The literature review revealed that the most important factors to influence the arsenic concentrations are the pH, the redox potential and the ferrous iron concentrations. The pH will influence arsenic concentrations directly through deprotonation. The higher the pH, the more arsenic is mobilised, which is visible in figure 8 of the results (Anawar et al., 2003; Ahmed et al., 2004; Nickson et al., 2000). The pH also influences the arsenic concentrations indirectly through the redox potential and the concentration of ferrous iron in the groundwater, which can be explained by the reduction hypothesis. A higher pH decreases the redox potential, which will transfer more ferric iron into ferrous iron, because of the more reduced environment (Sondergaard, 2009). Ferrous iron is more soluble and therefore has less adsorption sites, which mobilises arsenic (Sondergaard, 2009; Nickson et al., 2000; Van der Perk, 2013; Anawar et al., 2003), which is visible in figure 9 and 10. The molarity of bicarbonate, CO₂ and H₃O⁺ were also important, since these influence the pH through the equilibrium reaction of:



The data analysis found applicable formulas for the relations between the pH, redox potential, ferrous iron and arsenic concentrations and showed that the pH both has a direct and indirect effect on the arsenic concentrations as was described above. A multiple regression analysis showed that the pH is the most important variable to influence the arsenic concentrations in the Ganges delta, followed by the redox potential and ferrous iron (table 12, appendix 4). This can be expected due to the direct and indirect effect that the pH has on the arsenic concentrations.

After looking into the factors that influence the arsenic concentrations it should be made clear how much the effective sea level rise will amount to in the Ganges area. Therefore sub question 2 was: *How much will the effective sea level rise be in the Ganges delta from the year 2000 until the year 2100?* The effective sea level rise depends on three different factors, which are subsidence, tectonic uplift and the eustatic sea level rise. The numbers found for these factors are constants, so every year the increase or decrease is equal to the previous year. This explains why the effective sea level rises changes by the same amount for every year between 2000 and 2100. When combining these factors in Excel, the sea level rise will amount to 0.35 by the year 2100 and increases by 0.0035 per year (Pethick & Orford, 2013; Karim & Mimura, 2008; Shamsuddoha & Chowdhury, 2007; Paul & Rashid, 2017).

These previous two sub questions together lead to sub question 3: *How will the rising sea level change these factors and eventually thus the arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta?* The rising sea level will first increase the amount of bicarbonate ions and CO₂ ions in the groundwater, since diffusion will take place due to the molarity differences of bicarbonate and CO₂ (Van der Perk, 2013). More bicarbonate ions and CO₂ ions will be added to the groundwater due to the rising sea level, which will change the HCO₃⁻ molarity of the groundwater as can be seen in this equilibrium reaction:



Additionally, the HCO_3^- molarity in the groundwater changes even more due to the fact that HCO_3^- is released when ferric iron is transferred into ferrous iron due to the more reduced conditions (Nickson et al., 1998; formula 14, appendix III). The changed molarities of CO_2 and bicarbonate decrease the molarity of H_3O^+ , which will increase the pH, since these are connected through formula 11 (appendix III). The pH changes the redox potential, ferrous iron and arsenic concentration through a chain of formulas. The pH and redox potential are negatively related, which is displayed by the Nernst equation:

$$E_h = E_0 + \frac{0,05916}{n} \log \left(\frac{\{A\}\{B\}}{\{C\}\{D\}} \right) - \frac{0,05916h}{n} pH \text{ (Bundschuh \& Sracek, 2011).}$$

The redox potential relates also negatively to the amount of ferrous iron, since an electron wants to be released more when the redox potential is lower, which causes the oxidation of the new species (Oregon State University, n.d.; Sondergaard, 2009). More ferric iron will thus be transferred into ferrous iron (Sondergaard, 2009), which mobilizes arsenic (Nickson et al., 2000). The rising sea level thus relates positively to the molarity of CO_2 , HCO_3^- , the pH and concentration of ferrous iron, but negatively to the redox potential and the molarity of H_3O^+ due to the relation between all of these variables.

Each of these factors thus change the arsenic concentrations when these factors decrease or increase. But what is the overall effect on the arsenic concentrations in Bangladesh when all these variables are taken into account? This is what the main research question: *How will the effective sea level rise affect arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta from the year 2000 until the year 2100?* answers. Figure 14 displays that when all these factors are taken into account, the arsenic concentrations will increase due to an effective sea level rise of 0.35 m between the years 2000 and 2100. The effective sea level rise increases the concentrations of arsenic in the groundwater through a chain of formulas, since all the factors involved are related to each other, as can be seen in the formulas in appendix III and figure 15 in appendix II. The increase will amount to 317 $\mu\text{g/L}$, but the more the sea level increases, the slower the increase, according to table 13 through 22 (appendix III). This is due to the diffusion process; seawater keeps invading the groundwater, but the effect of this phenomenon will decrease over time. This is because the molarities of bicarbonate and CO_2 in the groundwater and seawater will differ less over time, which means that there are less molarity differences to diffuse (Van der Perk, 2013). The total concentrations of arsenic in 2100 will amount to 1017 $\mu\text{g/L}$, which is incredibly high, considering the limits of the World Health Organization, 10 $\mu\text{g/L}$, and Bangladesh, 50 $\mu\text{g/L}$ (Smith et al., 2002).

Compare findings to other research – validation

The calculated effective sea level rise was 0.35 m from 2000 to 2100 in this research. Kay et al., (2015) concluded that the expected sea level rise from 2000 until 2100 will be between 0.3 and 0.6 m, which corresponds to 0.35 m. Pethick & Orford (2013) also found a relative sea level rise (without taking tidal range amplification into account) of 0.28 m to 0.88 m between 2000 and 2100, which also corresponds to the calculated effective sea level rise.

The pH in the relevant regions of the groundwater of the Ganges delta is between 6.7 and 7.1, so an average of 6.9 is taken for the estimated pH in the groundwater (Bhattacharya et al., 2002). When the pH is 6.9, the redox potential is +46.693, according to formula 12 (see appendix III).

However, the model found a starting value of 7.34 for the pH and a starting value of -11.72 mV for the redox potential. This redox value is between the +590 and -440 mV, which are the values found in the research of Bhattacharya et al., (2002), so this value seems correct. There is an explanation for the deviation in pH, which is that little information could be found on the starting value of the CO₂ molarity in the groundwater of Bangladesh. Eventually, the molarity of CO₂ in groundwater from a different area was used, so there is some uncertainty about the H₃O⁺ concentration and thus the pH.

The starting concentration of ferrous iron in the groundwater found in the literature is 4.55 mg/L, since this is the average of data of multiple areas in the Ganges delta that are within the four coastal divisions. These are the areas of Satkhira and Laxmipur (Bhattacharya et al., 2002). However, this value differs from the value found in the model, which is 9.54 mg/L. An explanation is that only two of the four divisions were represented in the literature calculation, which causes the literature calculated value to be lower than the model value. There was also already some uncertainty about the amount of CO₂ in the groundwater, which influenced the pH and H₃O⁺ concentration, which again, due to the linear relation, can explain the deviation for ferrous iron.

In several articles the concentrations of arsenic are displayed, but these vary considerably. Again the distance to the coast is considered to be between 0 km and 80 km, so the areas within this distance are considered in the calculation (appendix I). In Aggerwal et al. (2000), the concentration of arsenic was 175 µg/L. Bhattacharya et al. (2002) calculated an arsenic concentration of 147 µg/l. Anawar et al. (2003) displayed an arsenic concentration of 317.27 µg/L. Anawar et al. (2002) calculated a much higher arsenic concentration of 703 µg/L and Tareq et al. (2003) calculated an arsenic concentration of 750 µg/L. Because these values vary considerably, the average value will be taken from all these sources, which is 418 µg/L. The calculated starting value for arsenic by the model is 701 µg/L. A potential reason for the lower value found in the literature is that the model uses one source to determine the molarity of bicarbonate and CO₂, since this source was the only source that had the full amount of information available. Meanwhile the literature-found concentration of arsenic used multiple sources to determine the average, which could lead to a different concentration. Also other factors do influence arsenic slightly, such as competition with phosphorus and the lowering of the groundwater table, which could lead to a lower concentration of arsenic when these factors would be included in the model. Another potential reason is again the uncertainty about the CO₂ molarity, which causes a higher pH, which due to the chain of events will overestimate the starting concentration of arsenic.

Limitations and strengths

This research has certain limitations. A limitation is that both iron and HCO₃⁻ act non-conservatively in reality, which means that they are taking part in other reactions and precipitate. In reality, a consistent relation between iron and arsenic and iron and bicarbonate is thus impossible, however in this research it has to be assumed (Mahanta et al., 2015). Also the sediments of the aquifer/groundwater were not taken into account. For a clear and full picture of the influence of sea level rise on the arsenic concentrations in the groundwater, this would have been useful. However, due to the short timeframe of this research, this was simply impossible. Lastly, the data was extrapolated into the future, while in the future certain

relationships between variables may change, for example due to climate change. This was an inevitable limitation.

This research however also has many strengths. Firstly, data is gathered from different articles, so that data is compared, which shows whether the chosen data is realistic. However, for some data, comparison is unfortunately impossible. Another strength is that this research can easily be replicated, since all the data and formulas are displayed. The results of this study are also compared to other studies, so that the research is validated.

Implications, further research and recommendations

This research gives insight into the increase of arsenic concentrations due to the rising sea level and hereby increased salt water intrusion. The implication is that the knowledge gap between sea level rise and the hereby changing arsenic concentrations is closed and long-term predictions about arsenic concentrations can be done. Because of this, the focus can move from short term solutions to more long term- and sustainable solutions to decrease the arsenic concentrations, since most solutions are currently short term (Smith et al., 2000). Further research can thus focus on making a long term plan to tackle the increasing arsenic concentrations, instead of focusing on short term solutions. Examples of long term solutions are using the deeper groundwater as drinking water, since this is arsenic-free water or implement better arsenic removal systems (Smith et al., 2000).

This research leads to the recommendation of implementing more measures by the government to reduce arsenic concentrations in the groundwater, especially in the coastal areas where the arsenic concentrations will increase most. However, this implies that the population is completely dependent on the government and cannot take action for their own. Therefore, there should also be solutions for the population of Bangladesh, such as the SONO filter, which has iron pores that filters the arsenic out of the groundwater. Unfortunately, often these measures are too expensive (Ahmed, 2001). It is recommended to make these measures cheaper, so that the population can afford them. This way they are not solely dependent on the government to ensure their health, but they can also fend for themselves (Smith et al., 2002).

Conclusion

In this thesis, the research question *How will the effective sea level rise affect arsenic concentrations in the groundwater of the four coastal divisions of the Ganges delta from the year 2000 until the year 2100?* was researched. This is of importance, since there is a knowledge gap between the sea level rise and arsenic concentrations, even though the population is heavily affected by both. Based on the literature review, the model in Excel and use of SPSS, several conclusions could be drawn. The pH influences the arsenic concentrations directly and indirectly through its relation to the redox potential and ferrous iron. The sea level will rise by 0.35 m between 2000 and 2100, which increases the molarity of bicarbonate and CO₂ by 2.5 mol/L and 0.03 mol/L. These changes will decrease the H₃O⁺ molarity by 2.8×10^{-8} mol/L, which increases the pH by 0.4. The pH influences the redox potential, which decreases by 55 mV, which again influences the ferrous iron concentration, which increases by 4.5 mg/L. The direct and indirect influence of the pH together change the arsenic concentrations in the groundwater of the four coastal divisions by 317 µg/L between 2000 and 2100 and this will reach a concentration of 1017 µg/L in 2100. Because of this sharp increase, it is recommended that more measures are implemented by the government to reduce the arsenic concentrations and that these measures are long-term, so that these measures are sustainable. Additionally, it must be ensured that the population can fend for themselves when it comes to the high arsenic concentrations, so that they are not completely dependent on the government.

References

- Aggarwal, P. K., Froehlich, K., Basu, A. R., Poreda, R. J., Kulkarni, K. M., Tarafdar, S. A., ... & Syed, R. A. (2000). *A report on isotope hydrology of groundwater in Bangladesh: implications for characterization and mitigation of arsenic in groundwater* (No. INIS-XA--648). International Atomic Energy Agency. Retrieved May 15, 2019 from <https://inis.iaea.org/collection/NCLCollectionStore/Public/34/065/34065511.pdf>
- Ahmed, K. M., Bhattacharya, P., Hasan, M. A., Akhter, S. H., Alam, S. M., Bhuyian, M. H., ... & Sracek, O. (2004). Arsenic enrichment in groundwater of the alluvial aquifers in Bangladesh: an overview. *Applied Geochemistry*, 19(2), 181-200.
- Ahmed, M. F. (2001, May). An overview of arsenic removal technologies in Bangladesh and India. In Proceedings of BUET-UNU international workshop on technologies for arsenic removal from drinking water, Dhaka (pp. 5-7).
- Anawar, H. M., Akai, J., Komaki, K., Terao, H., Yoshioka, T., Ishizuka, T., ... & Kato, K. (2003). Geochemical occurrence of arsenic in groundwater of Bangladesh: sources and mobilization processes. *Journal of Geochemical Exploration*, 77(2-3), 109-131.
- Anawar, H. M., Akai, J., Mostofa, K. M. G., Safiullah, S., & Tareq, S. M. (2002). Arsenic poisoning in groundwater: health risk and geochemical sources in Bangladesh. *Environment International*, 27(7), 597-604.
- Baier, T., & Neuwirth, E. (2003). High-level interface between R and Excel. Hornik et al.(2003), 68.
- Bangladesh on the globe. (June 22, 2014). Retrieved June 19, 2019 from [https://pt.m.wikipedia.org/wiki/Ficheiro:Bangladesh_on_the_globe_\(Bangladesh_centered\).svg](https://pt.m.wikipedia.org/wiki/Ficheiro:Bangladesh_on_the_globe_(Bangladesh_centered).svg)
- Bhattacharya, P., Jacks, G., Ahmed, K. M., Routh, J., & Khan, A. A. (2002). Arsenic in groundwater of the Bengal Delta Plain aquifers in Bangladesh. *Bulletin of Environmental Contamination and Toxicology*, 69(4), 538-545.
- Billo, E. J. (2004). *Excel for chemists: a comprehensive guide*. John Wiley & Sons.
- Bozlee, B.J. & Janebo, M. (2008). A Simplified Model To Predict the Effect of Increasing Atmospheric CO₂ on Carbonate Chemistry in the Ocean. *Journal of chemical education*, 85(2), 213-217.
- Brammer, H. (2014). Bangladesh's dynamic coastal regions and sea-level rise. *Climate risk management*, 1, 51-62.
- Brammer, H., & Ravenscroft, P. (2009). Arsenic in groundwater: a threat to sustainable agriculture in South and South-east Asia. *Environment International*, 35(3), 647-654.
- Brown, S., & Nicholls, R. J. (2015). Subsidence and human influences in mega deltas: The case of the Ganges–Brahmaputra–Meghna. *Science of The Total Environment*, 527–528, 362–374.
- Bundschuh, J., & Sracek, O. (2011). Hydrogeochemistry principles for geochemical modeling. *Geochemical Modeling of Groundwater, Vadose and Geothermal Systems*, 1.

- Field, A. (2018). *Discovering statistics using IBM SPSS Statistics*. London, England: SAGE.
- Google. (n.d.). *Divisions of Bangladesh*. Retrieved June 22, 2019 from <https://www.google.nl/maps/search/divisions+of+bangladesh/@23.5885983,88.0982366,7z/data=!3m1!4b1>
- Groundwater Geochemistry. (n.d.). Retrieved May 7, 2019 from <https://web.viu.ca/earle/geol304/304g.pdf>
- Hem, J. D. (1972). Chemical factors that influence the availability of iron and manganese in aqueous systems. *Geological Society of America Bulletin*, 83(2), 443-450.
- Hendriks, M. (2010). *Introduction to physical hydrology*. New York, US: Oxford University Press.
- Jain, A., Raven, K.P. & Loeppert, R.H. (1999). Arsenite and Arsenate adsorption on ferrihydrite: surface charge reduction and net OH₋ release stoichiometry. *Environmental Scientific Technology*, 33(8), 1179-1184.
- Karim, M. F., & Mimura, N. (2008). Impacts of climate change and sea-level rise on cyclonic storm surge floods in Bangladesh. *Global Environmental Change*, 18(3), 490-500.
- Kay, S., Caesar, J., Wolf, J., Bricheno, L., Nicholls, R. J., Saiful Islam, A. K. M., et al. (2015). Modelling the increased frequency of extreme sea levels in the Ganges–Brahmaputra–Meghna delta due to sea level rise and other effects of climate change. *Environmental Science: Processes & Impacts*, 17(7), 1311–1322.
- Kvaternyuk, S., Kaminska, G., Musz-Pomorska, A., ... & Sobczuk, H. (2016). *Water supply and wastewater removal*. Lublin Poland: Lublin University of Technology.
- Large detailed physical map of Bangladesh. (n.d.). Retrieved May 7, 2019 from <https://www.mapsland.com/asia/bangladesh/large-detailed-physical-map-of-bangladesh>
- Mahanta, C., Choudhury, R., Basu, S., Hemani, R., Dutta, A., Barua, P. P., ... & Saikia, L. (2015). Preliminary assessment of arsenic distribution in Brahmaputra River basin of India based on examination of 56,180 public groundwater wells. In *Safe and sustainable use of arsenic-contaminated aquifers in the Gangetic Plain* (pp. 57-64). Springer, Cham.
- Millero, F. J., Feistel, R., Wright, D. G., & McDougall, T. J. (2008). The composition of Standard Seawater and the definition of the Reference-Composition Salinity Scale. *Deep Sea Research Part I: Oceanographic Research Papers*, 55(1), 50-72.
- Ministry of foreign affairs. (2018). *Climate Change profile Bangladesh*. Retrieved May 20, 2019 from <https://www.google.nl/url?sa=t&rct=j&q=&esrc=s&source=web&cd=4&ved=2ahUKEwj417ztYfiAhUOKewKHfGxBZUQFjADegQIAhAC&url=https%3A%2F%2Fwww.government.nl%2Fbinaries%2Fgovernment%2Fdocuments%2Fpublications%2F2019%2F02%2F05%2Fclimate-change-profiles%2FBangladesh.pdf&usq=AOvVaw1INXIXcbi7IKppm0S0H96i>
- Nickson, R., McArthur, J., Burgess, W., Ahmed, K. M., Ravenscroft, P., & Rahman, M. (1998). Arsenic poisoning of Bangladesh groundwater. *Nature*, 395(6700), 338.

- Nickson, R. T., McArthur, J. M., Ravenscroft, P., Burgess, W. G., & Ahmed, K. M. (2000). Mechanism of arsenic release to groundwater, Bangladesh and West Bengal. *Applied Geochemistry*, 15(4), 403-413.
- Nordstrom, D.K. (2002). Worldwide occurrences of arsenic in ground water. *Science*, 296, 2143-2144.
- Oregon State University. (n.d.) Redox potential: oxidation and reduction reactions. Retrieved May 9, 2019 from http://oregonstate.edu/instruct/fw456/lecture/pdf/redox_potential.pdf
- Paul, B.K. & Rashid, H. (2017). *Climatic hazards in Coastal Bangladesh*. Oxford, UK: Heinemann-Butterworth.
- Pethick, J., & Orford, J. D. (2013). Rapid rise in effective sea-level in southwest Bangladesh: its causes and contemporary rates. *Global and Planetary Change*, 111, 237-245.
- Shamsuddoha, M., & Chowdhury, R. K. (2007). Climate change impact and disaster vulnerabilities in the coastal areas of Bangladesh. COAST Trust, Dhaka.
- Smith, A. H., Lingas, E. O., & Rahman, M. (2000). Contamination of drinking-water by arsenic in Bangladesh: a public health emergency. *Bulletin of the World Health Organization*, 78, 1093-1103.
- Smith, A.H., Lopipero, P.A., Bates, M.N. & Steinmaus, C.M. (2002). Arsenic epidemiology and drinking water standards. *Science*, 296, 2145.
- Snyder, M., Taillefert, M., & Ruppel, C. (2004). Redox zonation at the saline-influenced boundaries of a permeable surficial aquifer: effects of physical forcing on the biogeochemical cycling of iron and manganese. *Journal of Hydrology*, 296(1-4), 164-178.
- Solutions to the diffusion equation. (February, 2006). Retrieved June 3, 2019 from https://ocw.mit.edu/courses/materials-science-and-engineering/3-205-thermodynamics-and-kinetics-of-materials-fall-2006/lecture-notes/lecture03_slides.pdf
- Sondergaard, M. (2009). Redox potential. *Organic compounds*, 852-859
- Tareq, S. M., Safiullah, S., Anawar, H. M., Rahman, M. M., & Ishizuka, T. (2003). Arsenic pollution in groundwater: a self-organizing complex geochemical process in the deltaic sedimentary environment, Bangladesh. *Science of the total environment*, 313(1-3), 213-226.
- Van der Perk, M. (2013). *Soil and water contamination*. (2e ed.) Leiden, the Netherlands: Taylor & Francis Ltd.
- Van der Vecht, J.R. & Gijben, E.J. (2016). *Samengevat: vwo scheikunde*. Amersfoort, The Netherlands: ThiemeMeulenhoff.
- Werner, A. D., & Simmons, C. T. (2009). Impact of sea-level rise on sea water intrusion in coastal aquifers. *Groundwater*, 47(2), 197-204.
- Xyla, A. G., Sulzberger, B., Luther III, G. W., Hering, J. G., Van Cappellen, P., & Stumm, W. (1992). Reductive dissolution of manganese (III, IV)(hydr) oxides by oxalate: The effect of pH and light. *Langmuir*, 8(1), 95-103.

Yang, C., Delgado-Alonso, J., Hovorka, S., Mickler, P., Trevino, R., & Phillips, S. (2014). Monitoring dissolved CO₂ in groundwater for CO₂ leakage detection in a shallow aquifer. *Energy Procedia*, 63, 4209-4215.

Zeebe, R.E. (2011). On the molecular diffusion coefficients of dissolved CO₂, HCO₃⁻ and CO₃²⁻ and their dependence on isotopic mass. *Geochimica et Cosmochimica Acta*, 75(9), 2483-2498.

Appendix I: Additional calculations and explanation formulas

Calculation of the study area

The Ganges delta increases in salinity with 0.99%/year, which means that the area that is affected by the sea level rise changes (Paul & Rashid, 2017). In 2000, this area had a size of 1020750 ha and when the increase of 0.99%/year is taken into account the area will have a size of 2733736 ha in 2100, which is equivalent to 27337 km² (Paul & Rashid, 2017; table 2).

In google maps, the coastal length of Bangladesh, is revealed to be equivalent to a length of 342.50 km. This implicates that the salinity border lies 80 km inland in the year 2100 (27337/342.50). The divisions of Bangladesh that fall in this 80 km range are Khulna, Barisal, Comilla and Chittagong (Google, n.d.). These four coastal divisions determine the data that is taken into account and serve as a selection criteria for the model.

Table 2: The table shows how far the salinity will move inland due to the rising sea level.

SALINITY DEPTH INLANDS IN HA									
YEAR	Area (ha)	Year2	Area (ha)2	Year3	Area (ha)3	Year4	Area (ha)4	Year5	Area (ha)5
2000	1020750	2021	1255351	2042	1543871	2063	1898702	2084	2335085
2001	1030855	2022	1267779	2043	1559155	2064	1917499	2085	2358202
2002	1041060	2023	1280330	2044	1574591	2065	1936482	2086	2381549
2003	1051367	2024	1293005	2045	1590179	2066	1955654	2087	2405126
2004	1061775	2025	1305806	2046	1605922	2067	1975015	2088	2428937
2005	1072288	2026	1318734	2047	1621821	2068	1994567	2089	2452983
2006	1082903	2027	1331789	2048	1637877	2069	2014313	2090	2477268
2007	1093624	2028	1344974	2049	1654092	2070	2034255	2091	2501793
2008	1104451	2029	1358289	2050	1670467	2071	2054394	2092	2526560
2009	1115385	2030	1371736	2051	1687005	2072	2074733	2093	2551573
2010	1126427	2031	1385316	2052	1703706	2073	2095273	2094	2576834
2011	1137579	2032	1399031	2053	1720573	2074	2116016	2095	2602344,5
2012	1148841	2033	1412881	2054	1737607	2075	2136964	2096	2628108
2013	1160214	2034	1426869	2055	1754809	2076	2158120	2097	2654126
2014	1171700	2035	1440995	2056	1772182	2077	2179486	2098	2680402
2015	1183300	2036	1455261	2057	1789726	2078	2201063	2099	2706938
2016	1195015	2037	1469668	2058	1807445	2079	2222853	2100	2733736
2017	1206846	2038	1484217	2059	1825338	2080	2244859		
2018	1218793	2039	1498911	2060	1843409	2081	2267083		
2019	1230859	2040	1513750	2061	1861659	2082	2289528		
2020	1243045	2041	1528737	2062	1880089	2083	2312194		

Starting molarity bicarbonate

Aggarwal et al., (2000) have measured bicarbonate concentrations in multiple areas of the Ganges delta. There were 56 measurements at 56 locations of which 14 were inside the 80 km range. 3 of these 14 locations did not have any data, so the 11 locations that did have data on HCO₃⁻ were used to measure the average HCO₃⁻ concentration, which was 703 mg/L. This is similar to 0.703 g/l. To recalculate this value to the molarity, the atomic weight of HCO₃⁻ is needed, which is 61.01714 gram/mol (groundwater geochemistry, z.d.; Van der Vecht & Gijben, 2016). This gives a number of 0.703/61.01741=0.01152 mol/L. This value is used as the starting value of the bicarbonate molarity in groundwater.

Starting molarity CO₂

For CO₂, the background value in groundwater is 10⁻³ mol/L, so this value is used as a starting value in Excel for CO₂ in the groundwater (Yang et al., 2014).

Relation pH and redox potential

An increase of 1.5 pH, means a decrease of 200 mV. This relationship is displayed in formula 12 (see appendix III). The lower the redox potential, the faster reduction takes place, which means that there is an increase in ferrous iron and a decrease in ferric iron. This thus means that an increase in pH, which will likely happen due to the salt water intrusion, will result in an increase of soluble ferrous iron and thus arsenic concentrations (Sondergaard, 2009).

Relation between arsenic and iron hydroxide (Fe²⁺)

Bhattacharya et al. (2002) provide a graph that displays the linear relationship between the concentration of iron hydroxide and the concentration of arsenic, which is the first part of formula 15 (see appendix III): $y=36.53x + 40.28$. The higher the concentrations of ferrous iron, the higher the arsenic concentrations in the groundwater of the Ganges delta.

Relation between iron hydroxide (Fe²⁺) and redox potential

The relation between ferrous iron and the redox potential is represented by formula 13 (see appendix III) (Kvaternyuk et al., 2016). The lower the redox potential, the higher the concentration of ferrous iron in the groundwater.

Relationship between HCO₃⁻ and iron hydroxide

In the process of reduction, HCO₃⁻ is released into the water. This means that when the concentration of ferrous iron increases in the groundwater, the HCO₃⁻ concentration increases in the groundwater. This relationship is displayed by Nickson et al. (1998) and the formula belonging here is: $(\text{Fe(II)}) \cdot 25.734 - 9.7141$. This however is the formula for the concentration of HCO₃⁻ in mg/L and in the other formulas molarity is used. It thus needs to be in the unit of mol/L. Therefore the formula must be divided by 1000 and divided by the atomic weight of HCO₃⁻, which is 61.01714 g/mol (Van der Vecht & Gijben, 2016). The formula thus becomes $(\text{Fe(II)}) \cdot 25.734 - 9.7141 / 61017.14$, which is formula 14 in appendix III.

Relation pH and arsenic

The relation of pH and arsenic can be deduced from research that has both provided the pH and arsenic concentrations of the groundwater. When using the formula that is provided by Bhattacharya et al. (2002) the found formula is $y= 368.44x - 2391.8$, which is the second part of formula 15 (see appendix III).

Relation flux bicarbonate and CO₂

The formulas to calculate the flux of bicarbonate and CO₂ are formulas 4 and 5 (see appendix III). These fluxes represent the transfer of mol through a small area per time and these are the same for each year. When the fluxes are calculated it needs to be transferred to mol/L, so that the flux can be added to the current molarity in the groundwater. This is done by multiplying the flux by the amount of seconds in one year and dividing it by the area, which is the change in h each year (formulas 6 & 7, appendix III). This molarity that comes into the water as a flux needs to be added to the molarity that is already present in the groundwater (formulas 8 and 9, appendix III; Van der Perk, 2013; Solutions to the diffusion equation, 2006).

Appendix II: The model in Excel

Year	Effective se	Depth of fr	Depth of fr	Flux bic	Flux CO2	Molarity of flu	Molarity of flu	Molarity of flu	Molarity of flu	pH	Redox potential	When influence of Fe2+ on increase bicarbonate is taken into account										
												Fe2+	As concentrat	Molarity bic	Molarity bic	Molarity CO2	H2O+	pH	Redox potential	Fe2+	As concentrat	
2000	0	6	240	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.03645652	0.00130325	0.50355E-08	7.29640564	-6.15974642	9.02909898	668.931661	0.00367573	0.04014135	0.00130325	4.59077E-08	7.33811407	-11.7207487	9.54236509	700.712647
2001	0.0035	5.9965	239.86	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.06141124	0.0016065	3.699E-08	7.43191661	-24.2274147	10.5227751	772.188156	0.00294143	0.06507267	0.0016065	3.45739E-08	7.46125159	-28.1386739	10.8688049	794.504075
2002	0.007	5.993	239.72	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.08635686	0.00190976	3.127E-08	7.50489265	-33.9542173	11.3387006	872.777666	0.0064229	0.09099776	0.00190976	2.96813E-08	7.52751719	-36.9738666	11.5826884	845.03404
2003	0.0105	5.9895	239.58	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.11302048	0.00221301	2.8114E-08	7.55107211	-40.1144438	11.8364417	862.94217	0.00483282	0.1163353	0.00221301	2.69445E-08	7.56953152	-42.5765737	12.0353115	877.084124
2004	0.014	5.986	239.44	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.1362481	0.00251626	2.6114E-08	7.58312516	-44.3880769	12.1817566	887.40261	0.00497846	0.14122656	0.00251626	2.59355E-08	7.59871107	-46.4661473	12.3496647	899.282359
2005	0.0175	5.9825	239.3	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.16119372	0.00281951	2.4733E-08	7.60672436	-47.5345587	12.4599223	905.388323	0.00508568	0.1662794	0.00281951	2.39765E-08	7.62021469	-49.333225	12.5813246	915.676688
2006	0.021	5.979	239.16	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.18613994	0.00312277	2.3722E-08	7.62489437	-49.951401	12.6512521	919.199069	0.00518083	0.19130737	0.00312277	2.30811E-08	7.63674271	-51.5369054	12.759582	928.261707
2007	0.0245	5.9755	239.02	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.21108496	0.00342602	2.295E-08	7.63912105	-51.856943	12.786049	930.147869	0.00523332	0.21631827	0.00342602	2.23497E-08	7.64985398	-53.285031	12.9060605	938.352235
2008	0.028	5.972	238.88	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.23603508	0.00372927	2.234E-08	7.65089481	-53.439055	12.9118452	939.045238	0.00528637	0.24131695	0.00372927	2.18517E-08	7.66051437	-54.7063811	13.0154756	946.375238
2009	0.0315	5.9685	238.74	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.2609762	0.00403252	2.1849E-08	7.6605745	-54.7143975	13.0161233	946.410253	0.00530533	0.26630565	0.00403252	2.14114E-08	7.66955454	-55.851613	13.110721	953.111959
2010	0.035	5.965	238.6	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.28592182	0.00433578	2.1404E-08	7.66873097	-55.8019009	13.039986	952.636126	0.00536741	0.29128393	0.00433578	2.10471E-08	7.6768081	-56.878245	13.191009	958.790738
2011	0.0385	5.9615	238.46	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.3108674	0.00463903	2.1102E-08	7.67569872	-56.7309103	13.1790576	957.945409	0.00539907	0.31626605	0.00463903	2.07407E-08	7.68317669	-57.7279478	13.2596182	963.643471
2012	0.042	5.958	238.32	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.3358105	0.00494228	2.081E-08	7.68127068	-57.5338185	13.249325	962.534023	0.00542643	0.34123499	0.00494228	2.04794E-08	7.68882328	-58.4620223	13.3189314	967.83702
2013	0.0455	5.9545	238.18	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.36075867	0.00524553	2.056E-08	7.68697767	-58.2347323	13.3005664	966.539741	0.00545032	0.36620899	0.00524553	2.02454E-08	7.69348989	-59.1030605	13.3770229	971.501922
2014	0.049	5.951	238.04	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.38570429	0.00554879	2.0342E-08	7.69167005	-58.8519686	13.3504391	970.67242	0.00547135	0.39117564	0.00554879	2.00574E-08	7.69774299	-59.6675925	13.4163415	974.728527
2015	0.0525	5.9475	237.9	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.41064991	0.00585204	2.015E-08	7.69571513	-59.3996976	13.3946956	973.19751	0.00549001	0.41613993	0.00585204	1.98846E-08	7.70148278	-60.1668997	13.4568309	977.592346
2016	0.056	5.944	237.76	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.43559553	0.00615529	1.9981E-08	7.6993884	-59.890551	13.4342357	975.994185	0.00550669	0.44110222	0.00615529	1.97314E-08	7.70484123	-60.6184809	13.4930117	980.151418
2017	0.0595	5.9405	237.62	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.46054115	0.00645854	1.9836E-08	7.70268443	-60.3289151	13.4697763	978.507991	0.00552168	0.46606283	0.00645854	1.95947E-08	7.70786046	-61.0190348	13.525538	982.452011
2018	0.063	5.937	237.48	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.48548677	0.00676178	1.9694E-08	7.70565952	-60.7264369	13.5018961	980.779816	0.00553523	0.49101022	0.00676178	1.94725E-08	7.71058947	-61.3821893	13.5549378	984.531461
2019	0.0665	5.9335	237.34	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.51043239	0.00706505	1.9572E-08	7.70837367	-61.0874068	13.5310668	982.843055	0.00554753	0.51597992	0.00706505	1.93612E-08	7.71306824	-61.713885	13.5816418	986.420337
2020	0.07	5.93	237.2	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.53537801	0.0073683	1.9461E-08	7.71083476	-61.417988	13.5577373	984.752529	0.00555875	0.54093676	0.0073683	1.92605E-08	7.71532933	-62.0141929	13.6005	988.140477
2021	0.0735	5.9265	237.06	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.56032363	0.00767155	1.9359E-08	7.71310622	-61.7184517	13.5820509	986.441753	0.00556903	0.56589266	0.00767155	1.9169E-08	7.71713064	-62.2911206	13.6282325	989.71972
2022	0.077	5.923	236.92	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.58526925	0.00797481	1.9261E-08	7.71518168	-61.9957732	13.6044585	988.030674	0.00557848	0.59084773	0.00797481	1.90851E-08	7.71910364	-62.545075	13.6488421	991.173319
2023	0.0805	5.9195	236.78	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.61021487	0.00827806	1.9182E-08	7.71710409	-62.25198	13.625119	989.496089	0.0055872	0.61580207	0.00827806	1.90080E-08	7.72106326	-62.7793638	13.6677726	992.512279
2024	0.084	5.916	236.64	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.63516048	0.00858131	1.9104E-08	7.71888004	-62.4883288	13.644257	990.894016	0.00559527	0.64075576	0.00858131	1.8937E-08	7.72268948	-62.9961884	13.685292	993.751243
2025	0.0875	5.9125	236.5	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.66010611	0.00888456	1.9031E-08	7.72052836	-62.7080335	13.6620021	992.140625	0.00560275	0.66570886	0.00888456	1.88713E-08	7.72419858	-63.1974239	13.7015526	994.901541
2026	0.091	5.909	236.36	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.68505173	0.00918782	1.8961E-08	7.72206169	-62.9124849	13.6785288	993.203665	0.00560972	0.69066165	0.00918782	1.88103E-08	7.72560354	-63.3847197	13.7168854	995.971883
2027	0.0945	5.9055	236.22	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.70999735	0.00949107	1.8902E-08	7.72349223	-63.1032194	13.6939401	994.363111	0.00561622	0.71561357	0.00949107	1.87537E-08	7.72691407	-63.5594533	13.730808	996.970485
2028	0.098	5.902	236.08	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.73491297	0.00979432	1.8844E-08	7.72486992	-63.2815734	13.7083511	995.382403	0.0056223	0.74056257	0.00979432	1.87008E-08	7.72813962	-63.7228558	13.744009	997.904329
2029	0.1015	5.8985	235.94	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.75982859	0.01009757	1.8786E-08	7.72618352	-63.4487154	13.7226832	996.377618	0.00562799	0.76551658	0.01009757	1.86513E-08	7.7293282	-63.8758914	13.7558304	998.779723
2030	0.105	5.895	235.8	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.78474431	0.01040083	1.8739E-08	7.72726071	-63.6057093	13.7345382	997.34618	0.00563334	0.79046755	0.01040083	1.8601E-08	7.73053664	-64.0198102	13.768007	999.601421
2031	0.1085	5.8915	235.66	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.80965983	0.01070408	1.8691E-08	7.72816329	-63.753447	13.744702	998.27857	0.00563838	0.8154182	0.01070408	1.85517E-08	7.73138174	-64.1581273	13.7789343	1000.37476
2032	0.112	5.888	235.52	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.83457535	0.01100733	1.8646E-08	7.72912125	-63.8923553	13.7571169	999.187407	0.00564312	0.84036857	0.01100733	1.85029E-08	7.7323384	-64.2826785	13.7892404	1001.10371
2033	0.1155	5.8845	235.38	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.85949087	0.01131058	1.8604E-08	7.73007933	-64.0239555	13.7695355	999.625111	0.0056476	0.86531866	0.01131058	1.84549E-08	7.73321419	-64.4031141	13.7997916	1001.792
2034	0.119	5.881	235.24	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.88440639	0.01161384	1.8564E-08	7.73103741	-64.1482377	13.7837736	1000.3358	0.00565183	0.89026852	0.01161384	1.84061E-08	7.73409509	-64.5170337	13.8081747	1002.44294
2035	0.1225	5.8775	235.1	1.10743E-07	1.34625E-09	0.02494562	0.00003025	0.90932191	0.01191709	1.8526E-08	7.73199542	-64.2694838	13.7988886	1001.00031	0.00565584	0.91521815	0.01191709	1.83571E-08	7.73499059	-64.6248955	13.8	

Appendix III: Formulas and variables clarified

Table 3: The table displays all the independent variables in the model with their values, units and the sources where these were obtained.

Independent variables	Input	Unit	Source
Eustatic sea level rise	1.9	mm/year	Paul & Rashid, 2017
Subsidence	2.0	mm/year	Brammer, 2014
Tectonic uplift	3.6	mm/year	Paul & Rashid, 2017
Density fresh water	1.0	kg/L	Hendriks, 2010
Density seawater	1.025	kg/L	Hendriks, 2010
Molarity bicarbonate in seawater	0.00183	mol/L	Millero et al., 2008; Van der Vecht & Gijben, 2016
Diffusivity of bicarbonate	1.6×10^{-9}	m^2/s	Zeebe, 2011
Diffusivity of CO ₂	2.1×10^{-9}	m^2/s	Zeebe, 2011
Molarity CO ₂ in seawater	0.00001025	mol/L	Millero et al., 2008; Van der Vecht & Gijben, 2016
K _a	1.414×10^{-6}	No unit	Millero et al., 2008; Van der Vecht & Gijben, 2016

Table 4: The table displays all the dependent variables in the model with their starting values, units and the sources where these were obtained/when these were obtained through calculation by the model.

Dependent variables	Starting point	Unit	Source
Effective sea level rise	0	m	Pethick & Orford, 2013
Depth of fresh water above sea level	6	m	Tareq et al., 2003; Large detailed map of Bangladesh, n.d.
Depth of fresh water below sea level	240	m	Hendriks, 2010; Werner & Simmons, 2009
Flux CO ₂	1.34625×10^{-9}	$mol/m^2/s$	Van der Perk, 2013; Solutions to the diffusion equation, 2006
Flux bicarbonate	1.10743×10^{-7}	$mol/m^2/s$	Van der Perk, 2013; Solutions to the diffusion equation, 2006
Molarity of flux CO ₂	0.00030325	mol/L	Van der Perk, 2013; Solutions to the diffusion equation, 2006
Molarity of flux bicarbonate	0.02494562	mol/L	Van der Perk, 2013; Solutions to the diffusion equation, 2006
Molarity bicarbonate in aquifer	0.01152	mol/L	Groundwater geochemistry, n.d.; Van der Vecht & Gijben, 2016
Molarity CO ₂ in groundwater	0.001	mol/L	Yang et al., 2014
H ₃ O ⁺ in groundwater	4.59×10^{-8}	mol/L	Calculated by model
pH in groundwater	7.34	No unit	Calculated by model
Redox potential	-11.72	mV	Calculated by model
Fe ²⁺	9.54	mg/L	Calculated by model
Arsenic concentration	700.71	$\mu g/L$	Calculated by model

Table 5: The table displays all the formulas with their assigned number and the sources where these formulas were obtained.

	Formulas:	Sources for deriving formula
1	Effective Sea level Rise = Tectonic_uplift+Eustatic_sea_level_rise-Subsidence	Pethick & Orford, 2013; Karim & Mimura, 2008; Shamsuddoha & Chowdhury, 2007; Paul & Rashid, 2017; Brammer, 2014
2	Depth of fresh water above sea level= 6-Effective Sea level rise	Hendriks, 2010;
3	Depth of fresh water below sea level= (Density fresh water/Density seawater-Density freshwater)*Depth of fresh water above sea level	Hendriks, 2010; Werner & Simmons, 2009
4	Flux CO ₂ = $-D \cdot \frac{\Delta C}{\Delta x} \cdot 1000$	Van der Perk, 2013; Solutions to the diffusion equation, 2006; Zeebe, 2011
5	Flux Bicarbonate= $-D \cdot \frac{\Delta C}{\Delta x} \cdot 1000$	Van der Perk, 2013; Solutions to the diffusion equation, 2006; Zeebe, 2011
6	Molarity of flux CO ₂ = Flux CO ₂ *31536000*(1/change in depth of freshwater below sea level in 1 year)/1000	Van der Perk, 2013; Solutions to the diffusion equation, 2006.
7	Molarity of flux Bicarbonate= Flux bicarbonate*31536000*(1/change in depth of freshwater below sea level in 1 year)/1000	Van der Perk, 2013; Solutions to the diffusion equation, 2006.
8	Molarity of CO ₂ in groundwater after flux= Molarity CO ₂ previous year + molarity of flux CO ₂	Van der Perk, 2013; Solutions to the diffusion equation, 2006.
9	Molarity of bicarbonate in groundwater after flux= Molarity bicarbonate previous year + molarity of flux bicarbonate	Van der Perk, 2013; Solutions to the diffusion equation, 2006.
10	Molarity H ₃ O ⁺ = $K_a \cdot [\text{CO}_2] / [\text{HCO}_3^-]$	(Millero et al., 2008; Van der Vecht & Gijben, 2016).
11	pH= -LOG ₁₀ (H ₃ O ⁺)	Van der Vecht & Gijben, 2016
12	Redox potential= -133.33*pH+966.67	Sondergaard, 2009
13	Fe ²⁺ = -0.0808 *(redox potential)+8.5952	Kvaternyuk et al., 2016
14	HCO ₃ ⁻ = (Fe ²⁺ *25.734-9.7141)/61017.14	Van der Vecht & Gijben, 2016; Nickson et al., 1998

15	Arsenic concentration=	$36.53 \cdot \text{FeII} + 40.28 + 368.44 \cdot \text{pH} - 2391.8$	Bhattacharya et al., 2002
----	------------------------	---	---------------------------

Appendix III: Other graphs for data analysis derived from SPSS

Arsenic concentrations derived from the pH

Table 6: The table shows the significance and the Pearson correlation between the pH and the concentrations of arsenic.

		pH	As (µg/l)
pH	Pearson Correlation	1	,471**
	Sig. (1-tailed)		,003
	N	32	32
As (µg/l)	Pearson Correlation	,471**	1
	Sig. (1-tailed)	,003	
	N	32	32

** . Correlation is significant at the 0.01 level (1-tailed).

Table 7: The table shows the uncertainty/mean squared error of the correlation between pH and the concentrations of arsenic.

Tests of Between-Subjects Effects

Dependent Variable: As (µg/l)

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	592042,502 ^a	10	59204,250	1,262	,312	,375
Intercept	998085,170	1	998085,170	21,278	,000	,503
pH	592042,502	10	59204,250	1,262	,312	,375
Error	985066,034	21	46907,906			
Total	2872144,434	32				
Corrected Total	1577108,536	31				

a. R Squared = ,375 (Adjusted R Squared = ,078)

Arsenic concentrations derived from redox potential

Table 8: The table shows the significance and the Pearson correlation between the redox potential and the concentration of ferrous iron with and without bootstrapping.

		eh, mV	Fe (mg/l)
eh, mV	Pearson Correlation	1	-,477
	Sig. (1-tailed)		,116
	N	8	8
Bootstrap ^c	Bias	0	-,085

	Std. Error		0	,146
	95% Confidence Interval	Lower	1	-,904
		Upper	1	-,316
Fe (mg/l)	Pearson Correlation		-,477	1
	Sig. (1-tailed)		,116	
	N		8	8
	Bootstrap ^c	Bias	-,085	0
		Std. Error	,146	0
	95% Confidence Interval	Lower	-,904	1
		Upper	-,316	1

c. Unless otherwise noted, bootstrap results are based on 1000 bootstrap samples

Table 9: The table shows the uncertainty/mean squared error of the correlation between the redox potential and ferrous iron.

Tests of Between-Subjects Effects

Dependent Variable: Fe (mg/l)

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	290,674 ^a	5	58,135	17,280	,056	,977
Intercept	98,701	1	98,701	29,338	,032	,936
ehmV	290,674	5	58,135	17,280	,056	,977
Error	6,728	2	3,364			
Total	376,342	8				
Corrected Total	297,402	7				

a. R Squared = ,977 (Adjusted R Squared = ,921)

Arsenic concentrations derived from iron hydroxide

Table 10: The table shows the significance and the Pearson correlation between the concentrations of ferrous iron and arsenic.

Correlations

		Fe (mg/l)	As(µg/l)
Fe (mg/l)	Pearson Correlation	1	,645**
	Sig. (1-tailed)		,000
	N	33	33
As(µg/l)	Pearson Correlation	,645**	1
	Sig. (1-tailed)	,000	
	N	33	33

** . Correlation is significant at the 0.01 level (1-tailed).

Table 11: The table shows the uncertainty/mean squared error of the correlation between the concentrations of ferrous iron and arsenic.

Tests of Between-Subjects Effects

Dependent Variable: As($\mu\text{g/l}$)

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	1426788,379 ^a	28	50956,728	8,623	,024	,984
Intercept	1285547,120	1	1285547,120	217,544	,000	,982
F _{ell}	1426788,379	28	50956,728	8,623	,024	,984
Error	23637,500	4	5909,375			
Total	2730335,000	33				
Corrected Total	1450425,879	32				

a. R Squared = ,984 (Adjusted R Squared = ,870)

Multiple regression analysis

Table 12: This table shows which coefficients are most important to determine the arsenic concentrations. The pH has the highest influence, the redox potential has the middle influence and ferrous iron has the least influence, according to the standardised coefficients Beta.

Coefficients^a

Model		Unstandardized Coefficients		Standardized Coefficients	t	Sig.
		B	Std. Error	Beta		
1	(Constant)	522,778	,000		.	.
	eh, mV	,889	,000	,376	.	.
	pH	-72,222	,000	-,771	.	.
	Fe (mg/l)	-2,222	,000	-,151	.	.

a. Dependent Variable: As ($\mu\text{g/l}$)

Descriptive statistics

Table 13: The table shows the descriptive statistics of the arsenic concentrations from the year 2000 to 2009.

	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	700,712646688 92230	953,111958854 62280	879,828696992 465200	25,3411780444 22500	80,1358412122 26700	6421,753
Valid N (listwise)	10						

Table 14: The table shows the descriptive statistics of the arsenic concentrations from the year 2010 to 2019.

	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	958,790737563 92040	986,420237125 01690	974,765083192 895000	2,91512132748 2765	9,21842305057 9137	84,979
Valid N (listwise)	10						

Table 15: The table shows the descriptive statistics of the arsenic concentrations from the year 2020 to 2029.

	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	988,143447167 81850	998,779520764 92480	993,983020573 675300	1,12780789130 7412	3,56644169964 3038	12,720
Valid N (listwise)	10						

Table 16: The table shows the descriptive statistics of the arsenic concentrations from the year 2030 to 2039.

	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	999,601421446 64460	1005,23106000 627440	1002,61774324 3239500	,597850588853 307	1,89056956124 9323	3,574
Valid N (listwise)	10						

Table 17: The table shows the descriptive statistics of the arsenic concentrations from the year 2040 to 2049.

Descriptive Statistics – Year 2040 to 2049							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	1005,71062113 230620	1009,19449409 962680	1007,55079530 6703000	,370219570566 292	1,17073707735 8917	1,371
Valid N (listwise)	10						

Table 18: The table shows the descriptive statistics of the arsenic concentrations from the year 2050 to 2059.

Descriptive Statistics – Year 2050 to 2059							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	1009,5083715 0141850	1011,8767032 3309370	1010,7476402 81589400	,25176095345 7737	,79613803882 2093	,634
Valid N (listwise)	10						

Table 19: The table shows the descriptive statistics of the arsenic concentrations from the year 2060 to 2069.

Descriptive Statistics – Year 2060 to 2069							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	1012,09799873 747910	1013,81257151 020530	1012,98924265 7456500	,182301048162 305	,576486532028 937	,332
Valid N (listwise)	10						

Table 20: The table shows the descriptive statistics of the arsenic concentrations from the year 2070 to 2079.

Descriptive Statistics – Year 2070 to 2079							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	1013,97693970 521190	1015,27559097 610630	1014,64865580 2235700	,138096119195 321	,436698272687 313	,191
Valid N (listwise)	10						

Table 21: The table shows the descriptive statistics of the arsenic concentrations from the year 2080 to 2089.

Descriptive Statistics – Year 2080 to 2089							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	10	1013,97693970 521190	1015,27559097 610630	1014,64865580 2235700	,138096119195 321	,436698272687 313	,191
Valid N (listwise)	10						

Table 22: The table shows the descriptive statistics of the arsenic concentrations from the year 2090 to 2100.

Descriptive Statistics – Year 2090 to 2100							
	N	Minimum	Maximum	Mean		Std. Deviation	Variance
	Statistic	Statistic	Statistic	Statistic	Std. Error	Statistic	Statistic
As concentration	11	1016,52106631 425670	1017,42221728 382570	1016,98545749 8058200	,090084034648 977	,298774942532 031	,089
Valid N (listwise)	11						